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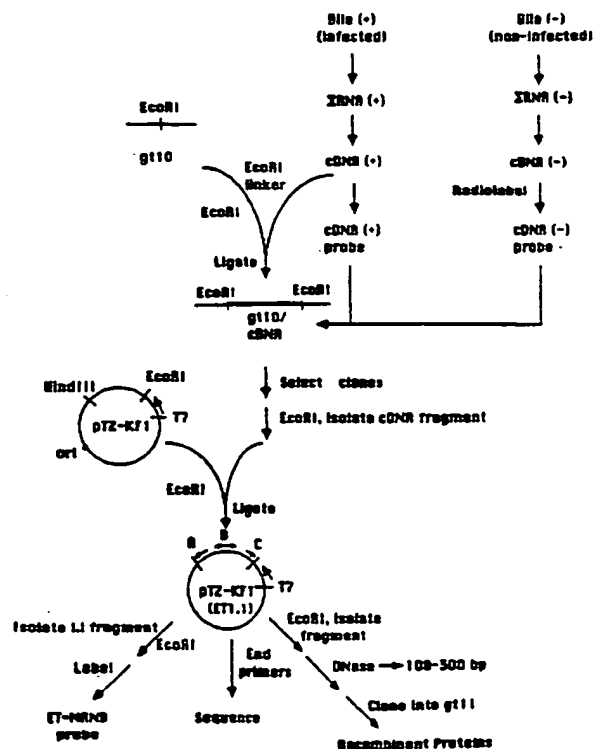
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(54) Title: ENTERICALLY TRANSMITTED NON-A/NON-B HEPATITIS VIRAL AGENT

(57) Abstract

Viral proteins derived from an enterically transmitted non-A/non-B viral hepatitis agent are disclosed. In one embodiment, the protein is immunologically reactive with antibodies present in individuals infected with the viral hepatitis agent. This protein is useful in a diagnostic method for detecting infection by the enterically transmitted agent. Also disclosed are DNA probes derived from a cloned sequence of the viral agent. These probes are useful for indentifying and sequencing the entire viral agent and for assaying the presence of the viral agent in an infected sample, using probe-specific amplification of virus-derived DNA fragments.



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ENTERERICALLY TRANSMITTED NON-A/NON-B
HEPATITIS VIRAL AGENT

5

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of
10 U.S. Application Serial No. 208,997, filed June 17,
1988, which is herein incorporated by reference.

INTRODUCTION

15 Field of Invention

This invention relates to recombinant
proteins, genes, and gene probes and more specifically
to such proteins and probes derived from an enterically
transmitted nonA/nonB hepatitis viral agent, and to
20 diagnostic methods and vaccine applications which
employ the proteins and probes.

Background

Enterically transmitted non-A/non-B (ET-NANB)
25 hepatitis viral agent is the reported cause of
hepatitis in several epidemics and sporadic cases in
Asia, Africa, and the Indian subcontinent. Infection
is usually by water contaminated with feces, although
the virus may also spread by close physical contact.
30 The virus does not seem to cause chronic infection.
The viral etiology in ET-NANB has been demonstrated by
infection of volunteers with pooled fecal isolates;
Immune electron microscopy (IEM) studies have shown
virus particles with 27-34 nm diameters in stools from
35 infected individuals. The virus particles reacted with
antibodies in serum from infected individuals from
geographically distinct regions, suggesting that a

single viral agent or class is responsible for the majority of ET-NANB hepatitis seen worldwide. No antibody reaction was seen in serum from individuals infected with blood-transmitted NANB virus, indicating
5 a different specificity between the two NANB types.

In addition to serological differences, the two types of NANB infection show distinct clinical differences. ET-NANB is characteristically an acute infection, often associated with fever and arthralgia,
10 and with portal inflammation and associated bile stasis in liver biopsy specimens (Arankalle). Symptoms are usually resolved within six weeks. Blood-transmitted NANB, by contrast, produces a chronic infection in about 50% of the cases. Fever and arthralgia are
15 rarely seen, and inflammation has a predominantly parenchymal distribution (Khuroo, 1980). The two viral agents can also be distinguished on the basis of primate host susceptibility. ET-NANB, but not the blood-transmitted agent, can be transmitted to
20 cynomolgus monkeys. The blood-transmitted agent is more readily transmitted to chimpanzees than is ET-NANB (Bradley, 1987).

There have been major efforts worldwide to identify and clone viral genomic sequences associated
25 with ET-NANB hepatitis. One goal of this effort, requiring virus-specific genomic sequences, is to identify and characterize the nature of the virus and its protein products. Another goal is to produce recombinant viral proteins which can be used in
30 antibody-based diagnostic procedures and for a vaccine. Despite these efforts, viral sequences associated with ET-NANB hepatitis have not been successfully identified or cloned heretofore, nor have
35 any virus-specific proteins been identified or produced.

Relevant Literature

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SUMMARY OF THE INVENTION

- Novel compositions, as well as methods of
25 preparation and use of the compositions are provided,
where the compositions comprise viral proteins and
fragments thereof derived from the viral agent for ET-
NANB. Methods for preparation of ET-NANB viral
proteins include isolating ET-NANB genomic sequences
30 which are then cloned and expressed in a host cell.
The resultant recombinant viral proteins find use as
diagnostic agents and as vaccines. The genomic
sequences and fragments thereof find use in preparing
ET-NANB viral proteins and as probes for virus
35 detection.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows vector constructions and manipulations used in obtaining and sequencing cloned ET-NANB fragment; and

5 Figures 2A-2B are representations of Southern blots in which a radiolabeled ET-NANB probe was hybridized with amplified cDNA fragments prepared from RNA isolated from infected (I) and non-infected (N) bile sources (2A), and from infected (I) and non-
10 infected (N) stool-sample sources (2B).

DESCRIPTION OF SPECIFIC EMBODIMENTS

Novel compositions comprising generic sequences and fragments thereof derived from the viral agent for ET-NANB are provided, together with
15 recombinant viral proteins produced using the genomic sequences and methods of using these compositions.

The genome of the ET-NANB viral agent is identified as containing a region which is homologous
20 to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1 (ET1.1) carried in E. coli strain BB4 and having ATCC deposit no. 67717. Initial studies sequenced the two terminal regions of the insert and an intermediate region. The 5'-end region of this insert contains the
25 sequence:

1		42
	GAT GGA AGG CAC TAA TCT GGC AAG ACC TGT CCC TGT TGC AGC	
30	43	84
	TGT TCT ACC ACC CTG CCC CGA GCT CGA ACA GGG CCT TCT CTA	
	85	126
	CCT GCC CCA GGA GCT CAC ACA CCC TGT GAT AGT GTC GTA ACA	
35	127	168
	TTT GAA TTA ACA GAC ATT GTG CAC TGC CGC ATG GCC GCC CCG	

169 210
AGC CAG CGC AAG GCC GTG CTG TCC ACA CTC GTG GGC CGC TAC

5 211
GGC.

An intermediate region has the sequence:

10 691 731
CTA GAG TGT GCT ATT ATG GAG GAG TGT GGG ATG CCG CAG TGG

733 774
CTC ATC CGC CTG TAT CAC CTT ATA AGG TCT GCG TGG ATC TTG

15 775 816
CAG GCC CCG AAG GAG TCT CTG CGA GGG TTT TGG AAG AAA CAC

817 858
20 TCC GGT GAG CCC GGC ACT CTT CTA TGG AAT ACT GTC TGG AAT

859 900
ATG GCC GTT ATT ACC CAC TGT TAT GAC TTC CGC GAT TTT CAG

25 901 942
GTG GCT GCC TTT AAA GGT GAT GAT TCG ATA GTG CTT TGC AGT

943 984
GAG TAT CGT CAG AGT CCA GGA GCT GCT GTC CTG ATC GCC GGC

30 985 1026
TGT GGC TTG AAG TTG AAG GTA GAT TTC CGC CCG ATC GGT TTG

1027
35 TAT.

The 3'-end region contains the sequence:

```

1191                                     1232
TGA GTA GAG GAT GTT GTT TCC CGT GTT TAT GGG GTT TCC CCT
5
1233                                     1274
GGA CTC GTT CAT AAC CTG ATT GGC ATG CTA CAG GCT GTT GCT

1275                                     1316
10 GAT GGC AAG GCA CAT TTC ACT GAG TCA GTA AAA CCA GTG CTC

1317      1327
GAC CGG AAT TC.

```

15 Additional work has provided the entire sequence, in both directions, as set forth below. The sequence of both strands is provided, since it is not known in which strand the gene is located. However, the sequence in one direction has been designated as the

20 "forward" sequence because of statistical similarities to known proteins. This sequence is set forth below along with the three possible translation sequences. There is one long open reading frame that starts at nucleotide 145 with an isoleucine and extends to the

25 end of the sequence. The two other reading frames have many termination codons. Standard abbreviations for nucleotides and amino acids are used here and elsewhere in this specification.

30 Forward Sequence

```

R I P P T D G R H Z S G K T C P C C S C
E F R Q L M E G T N L A R P V P V A A V
N S A N Z W K A L I W Q D L S L L Q L F
35      *      *      *      *      *
1      11      21      31      41      51
CGAATTCGCCCAACTGATGGAAGGCACTAATCTGGCAAGACCTGTCCCTGTTGCAGCTGT

```


S T T L P R A R T G P S L P A P G A H H
L P P C P E L E Q G L L Y L P Q E L T T
Y H P A P S S N R A F S T C P R S S P P
* * * * *
61 71 81 91 101 111
TCTACCACCCTGCCCCGAGCTCGAACAGGGCCTTCTCTACCTGCCCCAGGAGCTCACCAC

L Z Z C R N I Z I N R H C A L P H G R P
C D S V V T F E L T D I V H C R M A A P
V I V S Z H L N Z Q T L C T A A W P P R
* * * * *
121 131 141 151 161 171
CTGTGATAGTGTCTGAACATTTGAATTAACAGACATTGTGCACTGCCGCATGGCCGCCCC

E P A Q G R A V H T R G P L R R R T K L
S Q R K A V L S T L V G R Y G V A Q S S
A S A R P C C P H S W A A T A S H K A L
* * * * *
181 191 201 211 221 231
GAGCCAGCGCAAGGCCGTGCTGTCCACACTCGTGGGCGCTACGGCGTCGCACAAAGCTC

Y N A S H S D V R D S L A R F I P A I G
T M L P T L M F A T L S P V L S R P L A
Q C F P L Z C S R L S R P F Y P G H W P
* * * * *
241 251 261 271 281 291
TACAATGCTTCCCACTCTGATGTTGCGGACTCTCTCGCCCGTTTATCCCGGCCATTGGC

P V Q V T T C E L Y E L V E A M V E K G
P Y R L Q L V N C T S Z W R P W S R R A
R T G Y N L Z I V R A S G G H G R E G P
* * * * *
301 311 321 331 341 351
CCCGTACAGGTTACAACCTTGTAATTGTACGAGCTAGTGGAGGCCATGGTCGAGAAGGGC

Q D G S A V L E L D L C N R D V S R I T
R M A P P S L S L I F A T V T C P G S P
G W L R R P Z A Z S L Q P Z R V Q D H L
* * * * *
361 371 381 391 401 411
CAGGATGGCTCCGCCGTCCTTGAGCTTGATCTTTGCAACCGTGACGTGTCCAGGATCACC

F F Q K D C N K F T T G E T I A H G K V
S S R K I V T S S P Q V R P L P M V K W
L P E R L Z Q V H H R Z D H C P W Z S G
* * * * *
421 431 441 451 461 471
TTCTTCCAGAAAGATTGTAACAAGTTCACCACAGGTGAGACCATTGCCCATGGTAAAGTG

G Q G I S A W S K T F C A L F G P W F R
A R A S R P G A R P S A P S L A L G S A
P G H L G L E Q D L L R P L W P L V P R
* * * * *
481 491 501 511 521 531
GGCCAGGGCATCTCGGCCTGGAGCAAGACCTTCTGCGCCCTCTTTGGCCCTTGTTCCGC

A I E K A I L A L L P Q G V F Y G D A F
L L R R L F W P C S L R V C F T V M P L
Y Z E G Y S G P A P S G C V L R Z C L Z
* * * * *
541 551 561 571 581 591
GCTATTGAGAAGGCTATTCTGGCCCTGCTCCCTCAGGGTGTGTTTACGGTGATGCCTTT

D D T V F S A A V A A A K A S M V F E N
M T P S S R R L W P Q Q R H P W C L R M
Z H R L L G G C G R S K G I H G V Z E Z
* * * * *
601 611 621 631 641 651
GATGACACCGTCTTCTCGGCGGCTGTGGCCGAGCAAAGGCATCCATGGTGTGTTGAGAAT

D F S E F D S T Q N N F S L G L E C A I
T F L S L T P P R I T F L W V Z S V L L
L F Z V Z L H P E Z L F S G S R V C Y Y
* * * * *
661 671 681 691 701 711

GACTTTTCTGAGTTTGACTCCACCCAGAATAACTTTTCTCTGGGTCTAGAGTGTGCTATT

M E E C G M P Q W L I R L Y H L I R S A
W R S V G C R S G S S A C I T L Z G L R
G G V W D A A V A H P P V S P Y K V C V
* * * * *
721 731 741 751 761 771

ATGGAGGAGTGTGGGATGCCGAGTGGCTCATCCGCCTGTATCACCTTATAAGGTCTGCG

W I L Q A P K E S L R G F W K K H S G E
G S C R P R R S L C E G F G R N T P V S
D L A G P E G V S A R V L E E T L R Z A
* * * * *
781 791 801 811 821 831

TGGATCTTGCAGGCCCCGAAGGAGTCTCTGCGAGGGTTTGGGAAGAAACACTCCGGTGAG

P G T L L W N T V W N M A V I T H C Y D
P A L F Y G I L S G I W P L L P T V M T
R H S S M E Y C L E Y G R Y Y P L L Z L
* * * * *
841 851 861 871 881 891

CCCGGCACTCTTCTATGGAATACTGTCTGGAATATGGCCGTTATTACCCACTGTTATGAC

F R D F Q V A A F K G D D S I V L C S E
S A I F R W L P L K V M I R Z C F A V S
P R F S G G C L Z R Z Z F D S A L Q Z V
* * * * *
901 911 921 931 941 951

TTCCGCGATTTTCAGGTGGCTGCCTTTAAAGGTGATGATTCGATAGTGCTTTGCAGTGAG

Y R Q S P G A A V L I A G C G L K L K V
I V R V Q E L L S Z S P A V A Z S Z R Z
S S E S R S C C P D R R L W L E V E G R
* * * * *

961 971 981 991 1001 1011
TATCGTCAGAGTCCAGGAGCTGCTGTCCTGATCGCCGGCTGTGGCTTGAAGTTGAAGGTA

D F R P I G L Y A G V V V A P G L G A L
I S A R S V C M Q V L W W P P A L A R S
F P P D R F V C R C C G G P R P W R A P
* * * * *

1021 1031 1041 1051 1061 1071
GATTTCCGCCCCGATCGGTTTGTATGCAGGTGTTGTGGTGGCCCCCGGCCTTGGCGCGCTC

P D V V R F A G R L T E K N W G P G P E
L M L C A S P A G L P R R I G A L A L S
Z C C A L R R P A Y R E E L G P W P Z A
* * * * *

1081 1091 1101 1111 1121 1131
CCTGATGTTGTGCGCTTCGCCGGCCGGCTTACCGAGAAGAATTGGGGCCCTGGCCCTGAG

R A E Q L R L A V S D F L R K L T N V A
G R S S S A S L L V I S S A S S R M Z L
G G A A P P R C Z Z F P P Q A H E C S S
* * * * *

1141 1151 1161 1171 1181 1191
CGGGCGGAGCAGCTCCGCCTCGCTGTTAGTGATTTCCCTCCGCAAGCTCACGAATGTAGCT

Q M C V D V V S R V Y G V S P G L V H N
R C V W M L F P V F M G F P L D S F I T
D V C G C C F P C L W G F P W T R S Z P
* * * * *

1201 1211 1221 1231 1241 1251
CAGATGTGTGTGGATGTTGTTTCCCGTGTATGGGGTTTCCCTGGACTCGTTCATAAC

L I G M L Q A V A D G K A H F T E S V K
Z L A C Y R L L L M A R H I S L S Q Z N
D W H A T G C C Z W Q G T F H Z V S K T
* * * * *
1261 1271 1281 1291 1301 1311
CTGATTGGCATGCTACAGGCTGTTGCTGATGGCAAGGCACATTTCACTGAGTCAGTAAAA

P V L D R N S S
Q C S T G I R
S A R P E F E
* * *
1321 1331 1341
CCAGTGCTCGACCGGAATTCGAGC

The complimentary strand, referred to here as the "reverse sequence," is set forth below in the same manner as the forward sequence set forth above. Several open reading frames, shorter than the long open reading frame found in the forward sequence, can be seen in this reverse sequence.

Reverse Sequence

A R I P V E H W F Y Z L S E M C L A I S
L E F R S S T G F T D S V R C A L P S A
S N S G R A L V L L T Q Z N V P C H Q Q
* * * * *
1 11 21 31 41 51
GCTCGAATTCGGTCGAGCACTGGTTTTACTGACTCAGTGAAATGTGCCTTGCCATCAGC

N S L Z H A N Q V M N E S R G N P I N T
T A C S M P I R L Z T S P G E T P Z T R
Q P V A C Q S G Y E R V Q G K P H K H G
* * * * *
61 71 81 91 101 111
AACAGCCTGTAGCATGCCAATCAGGTTATGAACGAGTCCAGGGGAAACCCATAAACAGC

G N N I H T H L S Y I R E L A E E I T N
E T T S T H I Z A T F V S L R R K S L T
K Q H P H T S E L H S Z A C G G N H Z Q
* * * * *
121 131 141 151 161 171

GGAAACAACATCCACACACATCTGAGCTACATTCGTGAGCTTGCGGAGGAAATCACTAAC

S E A E L L R P L R A R A P I L L G K P
A R R S C S A R S G P G P Q F F S V S R
R G G A A P P A Q G Q G P N S S R Z A G
* * * * *
181 191 201 211 221 231

AGCGAGGCGGAGCTGCTCCGCGCGCTCAGGGCCAGGGCCCCAATTCTTCTCGGTAAGCCG

A G E A H N I R E R A K A G G H H N T C
P A K R T T S G S A P R P G A T T T P A
R R S A Q H Q G A R Q G R G P P Q H L H
* * * * *
241 251 261 271 281 291

GCCGCGGAAGCGCACAAACATCAGGGAGCGCGCCAAGGCCGGGGGCCACCACAACACCTGC

I Q T D R A E I Y L Q L Q A T A G D Q D
Y K P I G R K S T F N F K P Q P A I R T
T N R S G G N L P S T S S H S R R S G Q
* * * * *
301 311 321 331 341 351

ATACAAACCGATCGGGCGGAAATCTACCTTCAACTTCAAGCCACAGCCGGCGATCAGGAC

S S S W T L T I L T A K H Y R I I T F K
A A P G L Z R Y S L Q S T I E S S P L K
Q L L D S D D T H C K A L S N H H L Z R
* * * * *
361 371 381 391 401 411

AGCAGCTCCTGGACTCTGACGATACTCACTGCAAAGCACTATCGAATCATCACCTTTAAA

G S H L K I A E V I T V G N N G H I P D
A A T Z K S R K S Z Q W V I T A I F Q T
Q P P E N R G S H N S G Z Z R P Y S R Q
* * * * *

421 431 441 451 461 471
GGCAGCCACCTGAAAATCGCGGAAGTCATAACAGTGGGTAATAACGGCCATATTCCAGAC

S I P Z K S A G L T G V F L P K P S Q R
V F H R R V P G S P E C F F Q N P R R D
Y S I E E C R A H R S V S S K T L A E T
* * * * *

481 491 501 511 521 531
AGTATTCCATAGAAGAGTGCCGGGCTCACC GGAGTGTTCCTTCCAAAACCTCGCAGAGA

L L R G L Q D P R R P Y K V I Q A D E P
S F G A C K I H A D L I R Z Y R R M S H
P S G P A R S T Q T L Z G D T G G Z A T
* * * * *

541 551 561 571 581 591
CTCCTTCGGGGCCTGCAAGATCCACGCAGACCTTATAAGGTGATACAGGCGGATGAGCCA

L R H P T L L H N S T L Z T Q R K V I L
C G I P H S S I I A H S R P R E K L F W
A A S H T P P Z Z H T L D P E K S Y S G
* * * * *

601 611 621 631 641 651
CTGCGGCATCCCACTCCTCCATAATAGCACACTCTAGACCCAGAGAAAAGTTATTCTG

G G V K L R K V I L K H H G C L C C G H
V E S N S E K S F S N T M D A F A A A T
W S Q T Q K S H S Q T P W M P L L R P Q
* * * * *

661 671 681 691 701 711
GGTGGAGTCAAACCTCAGAAAAGTCATTCTCAAACACCATGGATGCCTTTGCTGCGGCCAC

S R R E D G V I K G I T V K H T L R E Q
A A E K T V S S K A S P Z N T P Z G S R
P P R R R C H Q R H H R K T H P E G A G
* * * * *

721 731 741 751 761 771
AGCCGCCGAGAAGACGGTGTTCATCAAAGGCATCACCGTAAAACACACCCTGAGGGAGCAG

G Q N S L L N S A E P R A K E G A E G L
A R I A F S I A R N Q G P K R A Q K V L
P E Z P S Q Z R G T K G Q R G R R R S C
* * * * *

781 791 801 811 821 831
GGCCAGAATAGCCTTCTCAATAGCGCGGAACCAAGGGCCAAAGAGGGCGCAGAAGGTCTT

A P G R D A L A H F T M G N G L T C G E
L Q A E M P W P T L P W A M V S P V V N
S R P R C P G P L Y H G Q W S H L W Z T
* * * * *

841 851 861 871 881 891
GCTCCAGGCCGAGATGCCCTGGCCCCACTTTACCATGGGCAATGGTCTCACCTGTGGTGAA

L V T I F L E E G D P G H V T V A K I K
L L Q S F W K K V I L D T S R L Q R S S
C Y N L S G R R Z S W T R H G C K D Q A
* * * * *

901 911 921 931 941 951
CTTGTTACAATCTTTCTGGAAGAAGGTGATCCTGGACACGTCACGGTTGCAAAGATCAAG

L K D G G A I L A L L D H G L H Z L V Q
S R T A E P S W P F S T M A S T S S Y N
Q G R R S H P G P S R P W P P L A R T I
* * * * *

961 971 981 991 1001 1011
CTCAAGGACGGCGGAGCCATCCTGGCCCTTCTCGACCATGGCCTCCACTAGCTCGTACAA

F T S C N L Y G A N G R D K T G E R V A
S Q V V T C T G P M A G I K R A R E S R
H K L Z P V R G Q W P G Z N G R E S R E
* * * * *

1021 1031 1041 1051 1061 1071
TTCACAAGTTGTAACCTGTACGGGGCCAATGGCCGGGATAAAACGGGCGAGAGAGTCGCG

N I R V G S I V E L C A T P Z R P T S V
T S E W E A L Z S F V R R R S G P R V W
H Q S G K H C R A L C D A V A A H E C G
* * * * *

1081 1091 1101 1111 1121 1131
AACATCAGAGTGGGAAGCATTGTAGAGCTTTGTGCGACGCCGTAGCGGCCACGAGTGTG

D S T A L R W L G A A M R Q C T M S V N
T A R P C A G S G R P C G S A Q C L L I
Q H G L A L A R G G H A A V H N V C Z F
* * * * *

1141 1151 1161 1171 1181 1191
GACAGCACGGCCTTGCGCTGGCTCGGGGCGGCCATGCGGCAGTGCACAATGTCTGTTAAT

S N V T T L S Q V V S S W G R Z R R P C
Q M L R H Y H R W Z A P G A G R E G P V
K C Y D T I T G G E L L G Q V E K A L F
* * * * *

1201 1211 1221 1231 1241 1251
TCAAATGTTACGACACTATCACAGGTGGTGAGCTCCTGGGGCAGGTAGAGAAGGCCCTGT

S S S G Q G G R T A A T G T G L A R L V
R A R G R V V E Q L Q Q G Q V L P D Z C
E L G A G W Z N S C N R D R S C Q I S A
* * * * *

1261 1271 1281 1291 1301 1311
TCGAGCTCGGGGACAGGTGGTAGAACAGCTGCAACAGGGACAGGTCTTGCCAGATTAGTG

P S I S W R N S
L P S V G G I
F H Q L A E F
* * *
1321 1331 1341
CCTTCCATCAGTTGGCGGAATTCG

DETAILED DESCRIPTION OF THE INVENTION

I. Definitions

The terms defined below have the following meaning herein:

1. "Enterically transmitted non-A/non-B (ET-NANB) hepatitis viral agent" means a virus, virus type, or virus class which (i) causes water-borne, infectious hepatitis, (ii) is transmissible in cynomolgus monkeys, (iii) is serologically distinct from hepatitis A virus (HAV) and hepatitis B virus (HAB), and (iv) includes a genomic region which is homologous to the 1.33 kb cDNA insert in plasmid pTZ-KP1(ET1.1) carried in E. coli strain BB4 identified by ATCC deposit number 67717.

2. Two double-strand nucleic acid fragments are "homologous" if their opposite strands are capable of hybridizing to one another under moderately stringent hybridization conditions, i.e., where hybridized strands contain at most about 5-10% basepair mismatches. A single-strand nucleic acid species is homologous to a double-strand fragment if it contains a region which is capable of hybridizing to one of the fragment strands under moderately stringent hybridization conditions.

3. A DNA fragment is "derived from" an ET-NANB viral agent if it has the same or substantially the

same basepair sequence as a region of the viral agent genome.

4. A protein is "derived from" an ET-NANB viral agent if it is encoded by an open reading frame of a DNA or RNA fragment derived from an ET-NANB viral agent.

II. Obtaining Cloned ET-NANB Fragments

- According to one aspect of the invention, it has been found that a virus-specific DNA clone can be produced by (a) isolating RNA from the bile of a cynomolgus monkey having a known ET-NANB infection, (b) cloning the cDNA fragments to form a fragment library, and (c) screening the library by differential hybridization to radiolabeled cDNAs from infected and non-infected bile sources.

A. cDNA Fragment Mixture

- ET-NANB infection in cynomolgus monkeys is initiated by inoculating the animals intravenously with a 10% w/v suspension from human case stools positive for 27-34 nm ET-NANB particles (mean diameter 32 nm). An infected animal is monitored for elevated levels of alanine aminotransferase, indicating hepatitis infection. ET-NANB infection is confirmed by immunospecific binding of seropositive antibodies to virus-like particles (VLPs), according to published methods (Gravelle). Briefly, a stool (or bile) specimen taken from the infected animal 3-4 weeks after infection is diluted 1:10 with phosphate-buffered saline, and the 10% suspension is clarified by low-speed centrifugation and filtration successively through 1.2 and 0.45 micron filters. The material may be further purified by pelleting through a 30% sucrose cushion (Bradley). The resulting preparation of VLPs is mixed with diluted serum from human patients with known ET-NANB infection. After incubation overnight,

the mixture is centrifuged overnight to pellet immune aggregates, and these are stained and examined by electron microscopy for antibody binding to the VLPs.

ET-NANB infection can also be confirmed by seroconversion to VLP-positive serum. Here the serum of the infected animal is mixed as above with 27-34 nm VLPs isolated from the stool specimens of infected human cases and examined by immune electron microscopy for antibody binding to the VLPs.

Bile can be collected from ET-NANB positive animals by either cannulating the bile duct and collecting the bile fluid or by draining the bile duct during necropsy. Total RNA is extracted from the bile by hot phenol extraction, as outlined in Example 1A. The RNA fragments are used to synthesize corresponding duplex cDNA fragments by random priming, also as referenced in Example 1A. The cDNA fragments may be fractionated by gel electrophoresis or density gradient centrifugation to obtain a desired size class of fragments, e.g., 500-4,000 basepair fragments.

Although alternative sources of viral material, such as VLPs obtained from stool samples (as described in Example 4), may be used for producing a cDNA fraction, the bile source is preferred. According to one aspect of the invention, it has been found that bile from ET-NANB-infected monkeys shows a greater number of intact viral particles than material obtained from stool samples, as evidenced by immune electron microscopy. Bile obtained from an ET-NANB infected human or cynomolgus monkey, for use as a source of ET-NANB viral protein or genomic material, or intact virus, forms part of the present invention.

B. cDNA Library and Screening

The cDNA fragments from above are cloned into a suitable cloning vector to form a cDNA library. This may be done by equipping blunt-ended fragments with a

suitable end linker, such as an EcoRI sequence, and inserting the fragments into a suitable insertion site of a cloning vector, such as at a unique EcoRI site. After initial cloning, the library may be recloned, if
5 desired, to increase the percentage of vectors containing a fragment insert. The library construction described in Example 1B is illustrative. Here cDNA fragments were blunt-ended, equipped with EcoRI ends, and inserted into the EcoRI site of the lambda phage
10 vector gtl0. The library phage, which showed less than 5% fragment inserts, was isolated, and the fragment inserts recloned into the lambda gtl0 vector, yielding more than 95% insert-containing phage.

The cDNA library is screened for sequences
15 specific for ET-NANB by differential hybridization to cDNA probes derived from infected and non-infected sources. cDNA fragments from infected and non-infected source bile or stool viral isolates can be prepared as above. Radiolabeling the fragments is by random
20 labeling, nick translation, or end labeling, according to conventional methods (Maniatis, p. 109). The cDNA library from above is screened by transfer to duplicate nitrocellulose filters, and hybridization with both infected-source and non-infected-source (control)
25 radiolabeled probes, as detailed in Example 2. Plaques which show selective hybridization to the infected-source probes are preferably replated at low plating density and rescreened as above, to isolate single clones which are specific for ET-NANB sequences. As
30 indicated in Example 2, sixteen clones which hybridized specifically with infected-source probes were identified by these procedures. One of the clones, designated lambda gtl0-1.1, contained a 1.33 kilobase fragment insert.

35

C. ET-NANB Sequences

The basepair sequence of cloned regions of the

ET-NANB fragments from Part B are determined by standard sequencing methods. In one illustrative method, described in Example 3, the fragment insert from the selected cloning vector is excised, isolated by gel electrophoresis, and inserted into a cloning vector whose basepair sequence on either side of the insertion site is known. The particular vector employed in Example 3 is a pTZ-KF1 vector shown at the left in Figure 1. The ET-NANB fragment from the gt10-1.1 phage was inserted at the unique EcoRI site of the pTZ-KF1 plasmid. Recombinants carrying the desired insert were identified by hybridization with the isolated 1.33 kilobase fragment, as described in Example 3. One selected plasmid, identified as pTZ-KF1 (ET1.1), gave the expected 1.33 kb fragment after vector digestion with EcoRI. E. coli strain BB4 infected with the pTZ-KF1(ET1.1) plasmid has been deposited with the American Type Culture Collection, Rockville, MD, and is identified by ATCC deposit number 67717.

The pTZ-KF1(ET1.1) plasmid is illustrated at the bottom in Figure 1. The fragment insert has 5' and 3' end regions denoted at A and C, respectively, and an intermediate region, denoted at B. The sequences in these regions were determined by standard dideoxy sequencing and are set forth above. The three short sequences (A, B, and C) are from the same insert strand. As will be seen in Example 3, the B-region sequence was actually determined from the opposite strand, so that the B-region sequence shown above represent the complement of the sequence in the sequenced strand. The base numbers of the partial sequences are approximate.

Later work in the laboratory of the inventors identified the full sequence, also set forth above. Fragments of this total sequence can readily be prepared using restriction endonucleases. Computer

analysis of both the forward and reverse sequence has identified a number of cleavage sites. The specific cleavage sites are summarized (for the forward direction) in the following tables.

5

	Pattern identifier (^ = cleavage site)	Pattern matched	Base number matched in "forward" strand			
	-----	-----	-----	-----	-----	-----
10	CC^WGG(BstNI)	CCWGG	106,	360,	410,	483,
			497,	973,	1129,	1243
	CC^SGG(NciI)	CCSGG	288,	841,	1063	
	GAAGANNNNNNN^ (MboII)	GAAGA	822,	1116		
15	TCTTC(<-7-MboII)	TCTTC	422,	611,	849	
	GCGTC(<-10-HgaI)	GCGTC	225			
	GCATCNNNNN^ (SfaI)	GCATC	488,	640		
	GCAGCNNNNNNN^ (BbvI)	GCAGC	53,	631,	1149	
	GCTGC(<-12-BbvI)	GCTGC	919,	979		
20	GGATGNNNNNNNN^ (FokI)	GGATG	363,	734,	1212	
	CATCC(<-13-FokI)	CATCC	641,	750		
	GGTGANNNNNNNN^ (HphI)	GGTGA	454,	589,	835,	931
	TCACC(<-7-HphI)	TCACC	114,	416,	446,	762
	GP^CGYC(AhaII)	GPCGYC	224			
25	GDGCH^C(BspI1286)	GDGCHC	77,	110,	158,	838,
			1125,	1324		
	GPGCY^C(BanII)	GPGCYC	77,	110,	838,	1125
	C^YCGPG(AvaI)	CYCGPG	74,	178		
	Y^GGCCP(EaeI)	YGGCCP	171,	290,	626,	875,
30			1101			
	GWGCW^C(GgiAI)	GWGCWC	77,	110,	158,	1324
	C^CTTGG(StyI)	CCTTGG	529,	1068		
	P^GATCY(XhoII)	PGATCY	782			
	CAG^CTG(PvuII)	CAGCTG	54			
35	C^CATGG(NcoI)	CCATGG	344,	468,	644	
	CGAT^CG(PvuI)	CGATCG	1031			
	C^GGCCG(EagI)	CGGCCG	1101			

	G [^] AATTC(EcoRI)	GAATTC	2, 1335
	GAGCT [^] C(SacI)	GAGCTC	77, 110
	GCATG [^] C(SphI)	GCATGC	1268
	GCC [^] GGC(NaeI)	GCCGGC	994, 1099, 1103
5	G [^] CGCGC(BssHII)	GCGCGC	1073
	GGGCC [^] C(ApaI)	GGGCCC	1125
	TCG [^] CGA(NruI)	TCGCGA	264
	T [^] CTAGA(XbaI)	TCTAGA	705
	TTT [^] AAA(DraI)	TTTAAA	925
10	G [^] TGCAC(ApaLI)	GTGCAC	158
	ACCTGCNNNN [^] (BspMI)	ACCTGC	99
	GCAGGT(<-8-BspMI)	GCAGGT	1045
	GACN [^] NNGTC(Tth111I)	GACNNNGTC	604
	CCANNNN [^] NTGG(PflMI)	CCANNNNNTGG	10
15	CC [^] TNAGG(MstII)	CCTNAGG	571
	GCCNNNN [^] NGGC(BglI)	GCCNNNNNGGC	216, 359, 738
	CCANNNNN [^] NTGG(BstXI)	CCANNNNNTGG	204

20 III. ET-NANB Fragments

According to another aspect, the invention includes ET-NANB-specific fragments or probes which hybridize with ET-NANB genomic sequences or cDNA fragments derived therefrom. The fragments may include

25 full-length cDNA fragments such as described in Section II, or may be derived from shorter sequence regions within cloned cDNA fragments. Shorter fragments can be prepared by enzymatic digestion of full-length

30 fragments under conditions which yield desired-sized fragments, as will be described in Section IV. Alternatively, the fragments can be produced by oligonucleotide synthetic methods, using sequences derived from the cDNA fragments. Methods or services for producing selected-sequence oligonucleotide

35 fragments are available.

To confirm that a given ET-NANB fragment is in fact derived from the ET-NANB viral agent, the fragment

can be shown to hybridize selectively with cDNA from infected sources. By way of illustration, to confirm that the 1.33 kb fragment in the pTZ-KF1(E1.1) plasmid is ET-NANB in origin, the fragment was excised from the pTZ-KF1(ET1.1) plasmid, purified, and radiolabeled by random labeling. The radiolabeled fragment was hybridized with fractionated cDNAs from infected and non-infected sources to confirm that the probe reacts only with infected-source cDNAs. This method is illustrated in Example 4, where the above radiolabeled 1.33 kb fragment from pTZ-KF1(ET1.1) plasmid was examined for binding to cDNAs prepared from infected and non-infected sources. The infected sources are (1) bile from a cynomolgus monkey infected with a strain of virus derived from stool samples from human patients from Burma with known ET-NANB infections and (2) a viral agent derived from the stool sample of a human ET-NANB patient from Mexico. The cDNAs in each fragment mixture were first amplified by a linker/primer amplification method described in Example 4. Fragment separation was on agarose gel, followed by Southern blotting and then hybridization to bind the radiolabeled 1.33 kb fragment to the fractionated cDNAs. The lane containing cDNAs from the infected sources showed a smeared band of bound probe, as expected (cDNAs amplified by the linker/primer amplification method would be expected to have a broad range of sizes). No probe binding to the amplified cDNAs from the non-infected sources was observed. The results indicate that the 1.33 kb probe is specific for cDNA fragments associated with ET-NANB infection. This same type of study, using ET 1.1 as the probe, has demonstrated hybridization to ET-NANB samples collected from Tashkent, Somalia, Borneo and Pakistan. Secondly, the fact that the probe is specific for ET-NANB related sequences derived from different continents (Asia, Africa and North America) indicates the cloned ET-NANB

Africa sequence is derived from a common ET-NANB virus or virus class responsible for ET-NANB hepatitis infection worldwide.

5 In a related confirmatory study, probe binding to fractionated genomic fragments prepared from human or cynomolgus monkey genomic DNA was examined. No probe binding was observed to either genomic fraction, demonstrating that the ET-NANB fragment is not an endogenous human or cynomologus fragment.

10 Another confirmation of ET-NANB specific sequences in the fragments is the ability to express ET-NANB proteins from coding regions in the fragments. Section IV below discusses methods of protein expression using the fragments.

15 One important use of the ET-NANB-specific fragments is for identifying ET-NANB-derived cDNAs which contain additional sequence information. The newly identified cDNAs, in turn, yield new fragment probes, allowing further iterations until the entire viral genome is identified and sequenced. Procedures for identifying additional ET-NANB library clones and generating new probes therefrom generally follow the cloning and selection procedures described in Section II.

25 The fragments (and oligonucleotides prepared based on the sequences given above) are also useful as primers for a polymerase chain reaction method of detecting ET-NANB viral genomic material in a patient sample. This diagnostic method will be described in Section V below.

30

IV. ET-NANB Proteins

35 As indicated above, ET-NANB proteins can be prepared by expressing open reading-frame coding regions in ET-NANB fragments. In one preferred approach, the ET-NANB fragments used for protein

expression are derived from cloned cDNAs which have been treated to produce desired-size fragments, and preferably random fragments with sizes predominantly between about 100 to about 300 base pairs. Example 5 describes the preparation of such fragments by DNAs digestion. Because it is desired to obtain peptide antigens of between about 30 to about 100 amino acids, the digest fragments are preferably size fractionated, for example by gel electrophoresis, to select those in the approximately 100-300 basepair size range.

A. Expression Vector

The ET-NANB fragments are inserted into a suitable expression vector. One exemplary expression vector is lambda gt11, which contains a unique EcoRI insertion site 53 base pairs upstream of the translation termination codon of the beta-galactosidase gene. Thus, the inserted sequence will be expressed as a beta-galactosidase fusion protein which contains the N-terminal portion of the beta-galactosidase gene, the heterologous peptide, and optionally the C-terminal region of the beta-galactosidase peptide (the C-terminal portion being expressed when the heterologous peptide coding sequence does not contain a translation termination codon). This vector also produces a temperature-sensitive repressor (c1857) which causes viral lysogeny at permissive temperatures, e.g., 32°C, and leads to viral lysis at elevated temperatures, e.g., 42°C. Advantages of this vector include: (1) highly efficient recombinant generation, (2) ability to select lysogenized host cells on the basis of host-cell growth at permissive, but not non-permissive, temperatures, and (3) high levels of recombinant fusion protein production. Further, since phage containing a heterologous insert produces an inactive beta-galactosidase enzyme, phage with inserts can be readily identified by a beta-galactosidase colored-substrate

reaction.

For insertion into the expression vector, the viral digest fragments may be modified, if needed, to contain selected restriction-site linkers, such as EcoRI linkers, according to conventional procedures. Example 1 illustrates methods for cloning the digest fragments into lambda gt11, which includes the steps of blunt-ending the fragments, ligating with EcoRI linkers, and introducing the fragments into EcoRI-cut lambda gt11. The resulting viral genomic library may be checked to confirm that a relatively large (representative) library has been produced. This can be done, in the case of the lambda gt11 vector, by infecting a suitable bacterial host, plating the bacteria, and examining the plaques for loss of beta-galactosidase activity. Using the procedures described in Example 1, about 50% of the plaques showed loss of enzyme activity.

20 B. Peptide Antigen Expression

The viral genomic library formed above is screened for production of peptide antigen (expressed as a fusion protein) which is immunoreactive with antiserum from ET-NANB seropositive individuals. In a preferred screening method, host cells infected with phage library vectors are plated, as above, and the plate is blotted with a nitrocellulose filter to transfer recombinant protein antigens produced by the cells onto the filter. The filter is then reacted with the ET-NANB antiserum, washed to remove unbound antibody, and reacted with reporter-labeled, anti-human antibody, which becomes bound to the filter, in sandwich fashion, through the anti-ET-NANB antibody.

Typically phage plaques which are identified by virtue of their production of recombinant antigen of interest are re-examined at a relatively low density for production of antibody-reactive fusion protein.

Several recombinant phage clones which produced immunoreactive recombinant antigen were identified in the procedure.

The selected expression vectors may be used for scale-up production, for purposes of recombinant protein purification. Scale-up production is carried out using one of a variety of reported methods for (a) lysogenizing a suitable host, such as E. coli, with a selected lambda gt11 recombinant (b) culturing the transduced cells under conditions that yield high levels of the heterologous peptide, and (c) purifying the recombinant antigen from the lysed cells.

In one preferred method involving the above lambda gt11 cloning vector, a high-producer E. coli host, BNN103, is infected with the selected library phage and replica plated on two plates. One of the plates is grown at 32°C, at which viral lysogeny can occur, and the other at 42°C, at which the infecting phage is in a lytic stage and therefore prevents cell growth. Cells which grow at the lower but not the higher temperature are therefore assumed to be successfully lysogenized.

The lysogenized host cells are then grown under liquid culture conditions which favor high production of the fused protein containing the viral insert, and lysed by rapid freezing to release the desired fusion protein.

C. Peptide Purification

The recombinant peptide can be purified by standard protein purification procedures which may include differential precipitation, molecular sieve chromatography, ion-exchange chromatography, isoelectric focusing, gel electrophoresis and affinity chromatography. In the case of a fused protein, such as the beta-galactosidase fused protein prepared as above, the protein isolation techniques which are used

can be adapted from those used in isolation of the native protein. Thus, for isolation of a beta-glactosidase fusion protein, the protein can be isolated readily by simple affinity chromatography, by
5 passing the cell lysis material over a solid support having surface-bound anti-beta-galactosidase antibody.

D. Viral Proteins

The ET-NANB protein of the invention may also
10 be derived directly from the ET-NANB viral agent. VLPs isolated from a stool sample from an infected individual, as above, are one suitable source of viral protein material. The VLPs isolated from the stool sample may be further purified by affinity
15 chromatography prior to protein isolation (see below). The viral agent may also be raised in cell culture, which provides a convenient and potentially concentrated source of viral protein. Co-owned U.S. Patent Application Serial No. 846,757, filed April 1,
20 1986, describes an immortalized trioma liver cell which supports NANB infection in cell culture. The trioma cell line is prepared by fusing human liver cells with a mouse/human fusion partner selected for human chromosome stability. Cells containing the desired
25 NANB viral agent can be identified by immunofluorescence methods, employing anti-ET-NANB human antibodies.

The viral agent is disrupted, prior to protein isolation, by conventional methods, which can include
30 sonication, high- or low-salt conditions, or use of detergents.

Purification of ET-NANB viral protein can be carried out by affinity chromatography, using a purified anti-ET-NANB antibody attached according to
35 standard methods to a suitable solid support. The antibody itself may be purified by affinity chromatography, where an immunoreactive recombinant ET-

NANB protein, such as described above, is attached to a solid support, for isolation of anti-ET-NANB antibodies from an immune serum source. The bound antibody is released from the support by standard methods.

- 5 Alternatively, the anti-ET-NANB antibody may be a monoclonal antibody (Mab) prepared by immunizing a mouse or other animal with recombinant ET-NANB protein, isolating lymphocytes from the animal and immortalizing the cells with a suitable fusion partner, and selecting
- 10 successful fusion products which react with the recombinant protein immunogen. These in turn may be used in affinity purification procedures, described above, to obtain native ET-NANB antigen.

15 V. Utility

 A. Diagnostic Methods

- The particles and antigens of the invention, as well as the genetic material, can be used in diagnostic
- 20 assays. Methods for detecting the presence of ET-NANB hepatitis comprise analyzing a biological sample such as a blood sample, stool sample or liver biopsy specimen for the presence of an analyte associated with ET-NANB hepatitis virus.

- 25 The analyte can be a nucleotide sequence which hybridizes with a probe comprising a sequence of at least about 16 consecutive nucleotides, usually 30 to 200 nucleotides, up to substantially the full sequence of the sequences shown above (cDNA sequences). The
- 30 analyte can be RNA or cDNA. The analyte is typically a virus particle suspected of being ET-NANB or a particle for which this classification is being ruled out. The virus particle can be further characterized as having an RNA viral genome comprising a sequence at least
- 35 about 80% homologous to a sequence of at least 12 consecutive nucleotides of the "forward" and "reverse" sequences given above, usually at least about 90%

homologous to at least about 60 consecutive nucleotides within the sequences, and may comprise a sequence substantially homologous to the full-length sequences. In order to detect an analyte, where the
5 analyte hybridizes to a probe, the probe may contain a detectable label.

The analyte can also comprise an antibody which recognizes an antigen, such as a cell surface antigen, on a ET-NANB virus particle. The analyte can also be a
10 ET-NANB viral antigen. Where the analyte is an antibody or an antigen, either a labelled antigen or antibody, respectively, can be used to bind to the analyte to form an immunological complex, which can then be detected by means of the label.

15 Typically, methods for detecting analytes such as surface antigens and/or whole particles are based on immunoassays. Immunoassays can be conducted either to determine the presence of antibodies in the host that have arisen from infection by ET-NANB hepatitis virus
20 or by assays that directly determine the presence of virus particles or antigens. Such techniques are well known and need not be described here in detail. Examples include both heterogeneous and homogeneous immunoassay techniques. Both techniques are based on the
25 formation of an immunological complex between the virus particle or its antigen and a corresponding specific antibody. Heterogeneous assays for viral antigens typically use a specific monoclonal or polyclonal antibody bound to a solid surface. Sandwich assays are
30 becoming increasingly popular. Homogeneous assays, which are carried out in solution without the presence of a solid phase, can also be used, for example by determining the difference in enzyme activity brought on by binding of free antibody to an enzyme-antigen
35 conjugate. A number of suitable assays are disclosed

in U.S. Patent Nos. 3,817,837, 4,006,360, 3,996,345.

When assaying for the presence of antibodies induced by ET-NANB viruses, the viruses and antigens of the invention can be used as specific binding agents to
5 detect either IgG or IgM antibodies. Since IgM antibodies are typically the first antibodies that appear during the course of an infection, when IgG synthesis may not yet have been initiated, specifically distinguishing between IgM and IgG antibodies present in the
10 blood stream of a host will enable a physician or other investigator to determine whether the infection is recent or chronic.

In one diagnostic configuration, test serum is reacted with a solid phase reagent having surface-bound
15 ET-NANB protein antigen. After binding anti-ET-NANB antibody to the reagent and removing unbound serum components by washing, the reagent is reacted with reporter-labeled anti-human antibody to bind reporter to the reagent in proportion to the amount of bound
20 anti-ET-NANB antibody on the solid support. The reagent is again washed to remove unbound labeled antibody, and the amount of reporter associated with the reagent is determined. Typically, the reporter is an enzyme which is detected by incubating the solid
25 phase in the presence of a suitable fluorometric or colorimetric substrate.

The solid surface reagent in the above assay prepared by known techniques for attaching protein material to solid support material, such as polymeric
30 beads, dip sticks, or filter material. These attachment methods generally include non-specific adsorption of the protein to the support or covalent attachment of the protein, typically through a free amine group, to a chemically reactive group on the
35 solid support, such as an activate carboxyl, hydroxyl, or aldehyde group.

In a second diagnostic configuration, known as

a homogeneous assay, antibody binding to a solid support produces some change in the reaction medium which can be directly detected in the medium. Known general types of homogeneous assays proposed heretofore
5 include (a) spin-labeled reporters, where antibody binding to the antigen is detected by a change in reported mobility (broadening of the spin splitting peaks), (b) fluorescent reporters, where binding is detected by a change in fluorescence efficiency, (c)
10 enzyme reporters, where antibody binding effects enzyme/substrate interactions, and (d) liposome-bound reporters, where binding leads to liposome lysis and release of encapsulated reporter. The adaptation of these methods to the protein antigen of the present
15 invention follows conventional methods for preparing homogeneous assay reagents.

In each of the assays described above, the assay method involves reacting the serum from a test individual with the protein antigen and examining the
20 antigen for the presence of bound antibody. The examining may involve attaching a labeled anti-human antibody to the antibody being examined, either IgM (acute phase) or IgG (convalescent phase), and measuring the amount of reporter bound to the solid
25 support, as in the first method, or may involve observing the effect of antibody binding on a homogeneous assay reagent, as in the second method.

Also forming part of the invention is an assay system or kit for carrying out the assay method just
30 described. The kit generally includes a support with surface-bound recombinant protein antigen which is (a) immunoreactive with antibodies present in individuals infected with enterically transmitted nonA/nonB viral agent and (b) derived from a viral hepatitis agent
35 whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. Coli strain BB4, and having

ATCC deposit no. 67717. A reporter-labeled anti-human antibody in the kit is used for detecting surface-bound anti-ET-NANB antibody.

5 B. Viral Genome Diagnostic Applications

 The genetic material of the invention can itself be used in numerous assays as probes for genetic material present in naturally occurring infections. One method for amplification of target nucleic acids, for later analysis by hybridization assays, is known as the polymerase chain reaction or PCR technique. The PCR technique can be applied to detecting virus particles of the invention in suspected pathological samples using oligonucleotide primers spaced apart from each other and based on the genetic sequence set forth above. The primers are complementary to opposite strands of a double stranded DNA molecule and are typically separated by from about 50 to 450 nt or more (usually not more than 2000 nt). This method entails preparing the specific oligonucleotide primers and then repeated cycles of target DNA denaturation, primer binding, and extension with a DNA polymerase to obtain DNA fragments of the expected length based on the primer spacing. Extension products generated from one primer serve as additional target sequences for the other primer. The degree of amplification of a target sequence is controlled by the number of cycles that are performed and is theoretically calculated by the simple formula 2^n where n is the number of cycles. Given that the average efficiency per cycle ranges from about 65% to 85%, 25 cycles produce from 0.3 to 4.8 million copies of the target sequence. The PCR method is described in a number of publications, including Saiki *et al.*, Science (1985) 230:1350-1354; Saiki *et al.*, Nature (1986) 324:163-166; and Scharf *et al.*, Science (1986) 233:1076-1078. Also see U.S. Patent Nos. 4,683,194; 4,683,195; and 4,683,202.

The invention includes a specific diagnostic method for determination of ET-NANB viral agent, based on selective amplification of ET-NANB fragments. This method employs a pair of single-strand primers derived from non-homologous regions of opposite strands of a DNA duplex fragment, which in turn is derived from an enterically transmitted viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4, and having ATCC deposit no. 67717. These "primer fragments," which form one aspect of the invention, are prepared from ET-NANB fragments such as described in Section III above. The method follows the process for amplifying selected nucleic acid sequences as disclosed in U.S. Patent No. 4,683,202, as discussed above.

C. Peptide Vaccine

Any of the antigens of the invention can be used in preparation of a vaccine. A preferred starting material for preparation of a vaccine is the particle antigen isolated from bile. The antigens are preferably initially recovered as intact particles as described above. However, it is also possible to prepare a suitable vaccine from particles isolated from other sources or non-particle recombinant antigens. When non-particle antigens are used (typically soluble antigens), proteins derived from the viral envelope or viral capsid are preferred for use in preparing vaccines. These proteins can be purified by affinity chromatography, also described above.

If the purified protein is not immunogenic per se, it can be bound to a carrier to make the protein immunogenic. Carriers include bovine serum albumin, keyhole limpet hemocyanin and the like. It is desirable, but not necessary, to purify antigens to be substantially free of human protein. However, it is more

important that the antigens be free of proteins, viruses, and other substances not of human origin that may have been introduced by way of, or contamination of, the nutrient medium, cell lines, tissues, or pathological fluids from which the virus is cultured or
5 obtained.

Vaccination can be conducted in conventional fashion. For example, the antigen, whether a viral particle or a protein, can be used in a suitable
10 diluent such as water, saline, buffered salines, complete or incomplete adjuvants, and the like. The immunogen is administered using standard techniques for antibody induction, such as by subcutaneous administration of physiologically compatible, sterile solutions
15 containing inactivated or attenuated virus particles or antigens. An immune response producing amount of virus particles is typically administered per vaccinating injection, typically in a volume of one milliliter or less.

20 A specific example of a vaccine composition includes, in a pharmacologically acceptable adjuvant, a recombinant protein or protein mixture derived from an enterically transmitted nonA/nonB viral hepatitis agent whose genome contains a region which is homologous to
25 the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4, and having ATCC deposit no. 67717. The vaccine is administered at periodic intervals until a significant titer of anti-ET-NANB antibody is detected in the serum. The vaccine
30 is intended to protect against ET-NANB infection.

D. Prophylactic and Therapeutic
Antibodies and Antisera

In addition to use as a vaccine, the compositions can be used to prepare antibodies to ET-NANB
35 virus particles. The antibodies can be used directly as antiviral agents. To prepare antibodies, a host

animal is immunized using the virus particles or, as appropriate, non-particle antigens native to the virus particle are bound to a carrier as described above for vaccines. The host serum or plasma is collected
5 following an appropriate time interval to provide a composition comprising antibodies reactive with the virus particle. The gamma globulin fraction or the IgG antibodies can be obtained, for example, by use of saturated ammonium sulfate or DEAE Sephadex, or other
10 techniques known to those skilled in the art. The antibodies are substantially free of many of the adverse side effects which may be associated with other anti-viral agents such as drugs.

The antibody compositions can be made even more
15 compatible with the host system by minimizing potential adverse immune system responses. This is accomplished by removing all or a portion of the Fc portion of a foreign species antibody or using an antibody of the same species as the host animal, for example, the use
20 of antibodies from human/human hybridomas.

The antibodies can also be used as a means of enhancing the immune response since antibody-virus complexes are recognized by macrophages. The anti-
bodies can be administered in amounts similar to those
25 used for other therapeutic administrations of antibody. For example, pooled gamma globulin is administered at 0.02-0.1 ml/lb body weight during the early incubation of other viral diseases such as rabies, measles and hepatitis B to interfere with viral entry
30 into cells. Thus, antibodies reactive with the ET-NANB virus particle can be passively administered alone or in conjunction with another anti-viral agent to a host infected with an ET-NANB virus to enhance the immune response and/or the effectiveness of an antiviral drug.

35 Alternatively, anti-ET-NANB-virus antibodies can be induced by administering anti-idiotypic antibodies as immunogens. Conveniently, a purified anti-

ET-NANB-virus antibody preparation prepared as described above is used to induce anti-idiotypic antibody in a host animal. The composition is administered to the host animal in a suitable diluent. Following administration, usually repeated administration, the host produces anti-idiotypic antibody. To eliminate an immunogenic response to the Fc region, antibodies produced by the same species as the host animal can be used or the Fc region of the administered antibodies can be removed. Following induction of anti-idiotypic antibody in the host animal, serum or plasma is removed to provide an antibody composition. The composition can be purified as described above for anti-ET-NANB-virus antibodies, or by affinity chromatography using anti-ET-NANB-virus antibodies bound to the affinity matrix. The anti-idiotypic antibodies produced are similar in conformation to the authentic ET-NANB antigen and may be used to prepare an ET-NANB vaccine rather than using a ET-NANB particle antigen.

When used as a means of inducing anti-ET-NANB-virus antibodies in a patient, the manner of injecting the antibody is the same as for vaccination purposes, namely intramuscularly, intraperitoneally, subcutaneously or the like in an effective concentration in a physiologically suitable diluent with or without adjuvant. One or more booster injections may be desirable. The anti-idiotypic method of induction of anti-ET-NANB-virus antibodies can alleviate problems which may be caused by passive administration of anti-ET-NANB-virus antibodies, such as an adverse immune response, and those associated with administration of purified blood components, such as infection with as yet undiscovered viruses.

The ET-NANB derived proteins of the invention are also intended for use in producing antiserum designed for pre- or post-exposure prophylaxis. Here an ET-NANB protein, or mixture of proteins is

formulated with a suitable adjuvant and administered by injection to human volunteers, according to known methods for producing human antisera. Antibody response to the injected proteins is monitored, during
5 a several- week period following immunization, by periodic serum sampling to detect the presence an anti-ET-NANB serum antibodies, as described in Section IIA above.

The antiserum from immunized individuals may be
10 administered as a pre-exposure prophylactic measure for individuals who are at risk of contracting infection. The antiserum is also useful in treating an individual post-exposure, analogous to the use of high titer antiserum against hepatitis B virus for post-exposure
15 prophylaxis.

E. Monoclonal Antibodies

For both in vivo use of antibodies to ET-NANB virus particles and proteins and anti-idiotypic anti-
20 bodies and diagnostic use, it may be preferable to use monoclonal antibodies. Monoclonal anti-virus particle antibodies or anti-idiotypic antibodies can be produced as follows. The spleen or lymphocytes from an immunized animal are removed and immortalized or used
25 to prepare hybridomas by methods known to those skilled in the art. To produce a human-human hybridoma, a human lymphocyte donor is selected. A donor known to be infected with a ET-NANB virus (where infection has been shown for example by the presence of anti-virus
30 antibodies in the blood or by virus culture) may serve as a suitable lymphocyte donor. Lymphocytes can be isolated from a peripheral blood sample or spleen cells may be used if the donor is subject to splenectomy. Epstein-Barr virus (EBV) can be used to immortalize
35 human lymphocytes or a human fusion partner can be used to produce human-human hybridomas. Primary in vitro immunization with peptides can also be used in the

generation of human monoclonal antibodies.

Antibodies secreted by the immortalized cells are screened to determine the clones that secrete antibodies of the desired specificity. For monoclonal
5 anti-virus particle antibodies, the antibodies must bind to ET-NANB virus particles. For monoclonal anti-idiotypic antibodies, the antibodies must bind to anti-virus particle antibodies. Cells producing antibodies of the desired specificity are selected.

10 The following examples illustrate various aspects of the invention, but are in no way intended to limit the scope thereof.

Material

15 The materials used in the following Examples were as follows:

Enzymes: DNase I and alkaline phosphatase were obtained from Boehringer Mannheim Biochemicals (BMB, Indianapolis, IN); EcoRI, EcoRI methylase, DNA ligase,
20 and DNA Polymerase I, from New England Biolabs (NEB, Beverly MA); and RNase A was obtained from Sigma (St. Louis, MO).

Other reagents: EcoRI linkers were obtained from NEB; and nitro blue tetrazolium (NBT),
25 5-bromo-4-chloro-3-indolyl phosphate (BCIP) 5-bromo-4-chloro-3-indolyl-B-D-galactopyranoside (X-gal) and isopropyl B-D-thiogalactopyranoside (IPTG) were obtained from Sigma.

30 cDNA synthesis kit and random priming labeling kits are available from Boehringer-Mannheim Biochemical (BMB, Indianapolis, IN).

Example 1

Preparing cDNA Library

35

A. Source of ET-NANB virus

Two cynomolgus monkeys (cynos) were

intravenously injected with a 10% suspension of a stool pool obtained from a second-passage cyno (cyno #37) infected with a strain of ET-NANB virus isolated from Burma cases whose stools were positive for ET-NANB, as evidenced by binding of 27-34 nm virus-like particles (VLPs) in the stool to immune serum from a known ET-NANB patient. The animals developed elevated levels of alanine aminotransferase (ALT) between 24-36 days after inoculation, and one excreted 27-34 nm VLPs in its bile in the pre-acute phase of infection.

The bile duct of each infected animal was cannulated and about 1-3 cc of bile was collected daily. RNA was extracted from one bile specimen (cyno #121) by hot phenol extraction, using a standard RNA isolation procedure. Double-strand cDNA was formed from the isolated RNA by a random primer for first-strand generation, using a cDNA synthesis kit obtained from Boehringer-Mannheim (Indianapolis, IN).

20 B. Cloning the Duplex Fragments

The duplex cDNA fragments were blunt-ended with T4 DNA polymerase under standard conditions (Maniatis, p. 118), then extracted with phenol/chloroform and precipitated with ethanol. The blunt-ended material was ligated with EcoRI linkers under standard conditions (Maniatis, pp. 396-397) and digested with EcoRI to remove redundant linker ends. Non-ligated linkers were removed by sequential isopropanol precipitation.

30 Lambda gt10 phage vector (Huynh) was obtained from Promega Biotec (Madison, WI). This cloning vector has a unique EcoRI cloning site in the phage cI repressor gene. The cDNA fragments from above were introduced into the EcoRI site by mixing 0.5 - 1.0 ug EcoRI-cleaved gt10, 0.5-3 µl of the above duplex fragments, 0.5 µl 10X ligation buffer, 0.5 µl ligase (200 units), and distilled water to 5 µl. The mixture

was incubated overnight at 14°C, followed by in vitro packaging, according to standard methods (Maniatis, pp. 256-268).

5 The packaged phage were used to infect an E. coli hfl strain, such as strain HG415. Alternatively, E. coli, strain C600 hfl, available from Promega Biotec, Madison, WI, could be used. The percentage of recombinant plaques obtained with insertion of the EcoRI-ended fragments was less than 5% by analysis of
10 20 random plaques.

 The resultant cDNA library was plated and phage were eluted from the selection plates by addition of elution buffer. After DNA extraction from the phage, the DNA was digested with EcoRI to release the
15 heterogeneous insert population, and the DNA fragments were fractionated on agarose to remove phage fragments. The 500-4,000 basepair inserts were isolated and recloned into lambda gt10 as above, and the packaged phage was used to infect E. coli strain
20 HG415. The percentage of successful recombinants was greater than 95%. The phage library was plated on E. coli strain HG415, at about 5,000 plaques/plate, on a total of 8 plates.

25

30

35

Example 2Selecting ET-NANB Cloned Fragments

A. cDNA Probes

Duplex cDNA fragments from noninfected and ET-
5 NANB-infected cynomolgus monkeys were prepared as in
Example 1. The cDNA fragments were radiolabeled by
random priming, using a random-priming labeling kit
obtained from Boehringer-Mannheim (Indianapolis, IN).

10 B. Clone Selection

The plated cDNA library from Example 1 was
transferred to each of two nitrocellulose filters, and
the phage DNA was fixed on the filters by baking,
according to standard methods (Maniatis, pp. 320-
15 323). The duplicate filters were hybridized with
either infected-source or control cDNA probes from
above. Autoradiographs of the filters were examined to
identify library clones which hybridized with
radiolabeled cDNA probes from infected source only,
20 i.e., did not hybridize with cDNA probes from the non-
infected source. Sixteen such clones, out of a total
of about 40,000 clones examined, were identified by
this subtraction selection method.

Each of the sixteen clones was picked and
25 replated at low concentration on an agar plate. The
clones on each plate were transferred to two nitro-
cellulose as duplicate lifts, and examined for hybrid-
ization to radiolabeled cDNA probes from infected and
noninfected sources, as above. Clones were selected
30 which showed selective binding for infected-source
probes (i.e., binding with infected-source probes and
substantially no binding with non-infected-source
probes). One of the clones which bound selectively to
probe from infected source was isolated for further
35 study. The selected vector was identified as lambda
gt10-1.1, indicated in Figure 1.

Example 3ET-NANB Sequence

Clone lambda gtl0-1.1 from Example 2 was digested with EcoRI to release the heterologous insert, which was separated from the vector fragments by gel electrophoresis. The electrophoretic mobility of the fragment was consistent with a 1.33 kb fragment. This fragment, which contained EcoRI ends, was inserted into the EcoRI site of a pTZ-KF1 vector, whose construction and properties are described in co-owned U.S. patent application for "Cloning Vector System and Method for Rare Clone Identification", Serial No. 125, 650, filed November 25, 1987. Briefly, and as illustrated in Figure 1, this plasmid contains a unique EcoRI site adjacent a T7 polymerase promoter site, and plasmid and phage origins of replication. The sequence immediately adjacent each side of the EcoRI site is known. E. coli BB4 bacteria, obtained from Stratagene (La Jolla, CA, were transformed with the plasmid.

Radiolabeled ET-NANB probe was prepared by excising the 1.33 kb insert from the lambda gtl0-1.1 phage in Example 2, separating the fragment by gel electrophoresis, and randomly labeling as above. Bacteria transfected with the above pTZ-KF1 and containing the desired ET-NANB insert were selected by replica lift and hybridization with the radiolabeled ET-NANB probe, according to methods outlined in Example 2.

One bacterial colony containing a successful recombinant was used for sequencing a portion of the 1.33 kb insert. This isolate, designated pTZ-KF1(ET1.1), has been deposited with the American Type Culture Collection, and is identified by ATCC deposit no. 67717. Using a standard dideoxy sequencing procedure, and primers for the sequences flanking the EcoRI site, about 200-250 basepairs of sequence from the 5'-end region and 3'-end region of the insert were

obtained. The sequences are given above in Section II. Later sequencing by the same techniques gave the full sequence in both directions, also given above.

5

Example 4Detecting ET-NANB Sequences

cdNA fragment mixtures from the bile of noninfected and ET-NANB-infected cynomolgus monkeys were prepared as above. The cdNA fragments obtained from human stool samples were prepared as follows. Thirty ml of a 10% stool suspension obtained from an individual from Mexico diagnosed as infected with ET-NANB as a result of an ET-NANB outbreak, and a similar volume of stool from a healthy, non-infected individual, were layered over a 30% sucrose density gradient cushion, and centrifuged at 25,000 xg for 6 hr in an SW27 rotor, at 15°C. The pelleted material from the infected-source stool contained 27-34 nm VLP particles characteristic of ET-NANB infection in the infected-stool sample. RNA was isolated from the sucrose-gradient pellets in both the infected and non-infected samples, and the isolated RNA was used to produce cdNA fragments as described in Example 1.

25

The cdNA fragment mixtures from infected and non-infected bile source, and from infected and non-infected human-stool source were each amplified by a novel linker/primer replication method described in co-owned patent application serial number 07/208,512 for "DNA Amplification and Subtraction Technique," filed June 17, 1988. Briefly, the fragments in each sample were blunt-ended with DNA Pol I then extracted with phenol/chloroform and precipitated with ethanol. The blunt-ended material was ligated with linkers having the following sequence:

30
35

5'-GGAATTCGCGCCGCTCG-3'
3'-TTCCTTAAGCGCCGCGAGC-5'

The duplex fragments were digested with NruI to remove linker dimers, mixed with a primer having the sequence 5'-GGAATTCGCGCCGCTCG-3', and then heat denatured and cooled to room temperature to form single-strand DNA/primer complexes. The complexes were replicated to form duplex fragments by addition of *Thermus aquaticus* (Taq) polymerase and all four deoxynucleotides. The replication procedures, involving successive strand denaturation, formation of strand/primer complexes, and replication, was repeated 25 times.

The amplified cDNA sequences were fractionated by agarose gel electrophoresis, using a 2% agarose matrix. After transfer of the DNA fragments from the agarose gels to nitrocellulose paper, the filters were hybridized to a random-labeled ³²P probe prepared by (i) treating the pTZ-KF1(ET1.1) plasmid from above with EcoRI, (ii) isolating the released 1.33 kb ET-NANB fragment, and (iii) randomly labeling the isolated fragment. The probe hybridization was performed by conventional Southern blotting methods (Maniatis, pp. 382-389). Figure 2 shows the hybridization pattern obtained with cDNAs from infected (I) and non-infected (N) bile sources (2A) and from infected (I) and non-infected (N) human stool sources (2B). As seen, the ET-NANB probe hybridized with fragments obtained from both of the infected sources, but was non-homologous to sequences obtained from either of the non-infected sources, thus confirming the specificity of derived sequence.

Southern blots of the radiolabeled 1.33 kb fragment with genomic DNA fragments from both human and cynomolgus-monkey DNA were also prepared. No probe hybridization to either of the genomic fragment

mixtures was observed, confirming that the ET-NANB sequence is exogenous to either human or cynomolgus genome.

5

Example 5

Expressing ET-NANB Proteins

A. Preparing ET-NANB Coding Sequences

The pTZ-KF1(ET1.1) plasmid from Example 2 was
10 digested with EcoRI to release the 1.33 kb ET-NANB
insert which was purified from the linearized plasmid
by gel electrophoresis. The purified fragment was
suspended in a standard digest buffer (0.5M Tris HCl,
pH 7.5; 1 mg/ml BSA; 10mM MnCl₂) to a concentration of
15 about 1 mg/ml and digested with DNase I at room
temperature for about 5 minutes. These reaction
conditions were determined from a prior calibration
study, in which the incubation time required to produce
predominantly 100-300 basepair fragments was
20 determined. The material was extracted with
phenol/chloroform before ethanol precipitation.

The fragments in the digest mixture were blunt-
ended and ligated with EcoRI linkers as in Example 1.
The resultant fragments were analyzed by
25 electrophoresis (5-10V/cm) on 1.2% agarose gel, using
PhiX174/HaeIII and lambda/HindIII size markers. The
100-300 bp fraction was eluted onto NA45 strips
(Schleicher and Schuell), which were then placed into
1.5 ml microtubes with eluting solution (1 M NaCl, 50
30 mM arginine, pH 9.0), and incubated at 67°C for 30-60
minutes. The eluted DNA was phenol/chloroform
extracted and then precipitated with two volumes of
ethanol. The pellet was resuspended in 20 µl TE (0.01
M Tris HCl, pH 7.5, 0.001 M EDTA).

35

B. Cloning in an Expression Vector

Lambda gt11 phage vector (Huynh) was obtained

from Promega Biotec (Madison, WI). This cloning vector has a unique EcoRI cloning site 53 base pairs upstream from the beta-galactosidase translation termination codon. The genomic fragments from above were
5 introduced into the EcoRI site by mixing 0.5-1.0 µg EcoRI-cleaved gtl1, 0.3-3 µl of the above sized fragments, 0.5 µl 10X ligation buffer (above), 0.5 µl ligase (200 units), and distilled water to 5 µl. The mixture was incubated overnight at 14°C, followed by in
10 vitro packaging, according to standard methods (Maniatis, pp. 256-268).

The packaged phage were used to infect E. coli strain KM392, obtained from Dr. Kevin Moore, DNAX (Palo Alto, CA). Alternatively, E. Coli strain Y1090,
15 available from the American Type Culture Collection (ATCC #37197), could be used. The infected bacteria were plated and the resultant colonies were checked for loss of beta-galactosidase activity-(clear plaques) in the presence of X-gal using a standard X-gal substrate
20 plaque assay method (Maniatis). About 50% of the phage plaques showed loss of beta-galactosidase enzyme activity (recombinants).

C. Screening for ET-NANB Recombinant Proteins

25 ET-NANB convalescent antiserum was obtained from patients infected during documented ET-NANB outbreaks in Mexico, Borneo, Pakistan, Somalia, and Burma. The sera were immunoreactive with VLPs in stool specimens from each of several other patients with ET-
30 NANB hepatitis.

A lawn of E. coli KM392 cells infected with about 10⁴ pfu of the phage stock from above was prepared on a 150 mm plate and incubated, inverted, for 5-8 hours at 37°C. The lawn was overlaid with a
35 nitrocellulose sheet, causing transfer of expressed ET-NANB recombinant protein from the plaques to the paper. The plate and filter were indexed for matching

corresponding plate and filter positions.

The filter was washed twice in TBST buffer (10 mM Tris, pH 8.0, 105 mM NaCl, 0.05% Tween 20), blocked with AIB (TBST buffer with 1% gelatin), washed again in
5 TBST, and incubated overnight after addition of antiserum (diluted to 1:50 in AIB, 12-15 ml/plate). The sheet was washed twice in TBST and then contacted with enzyme-labeled anti-human antibody to attach the labeled antibody at filter sites containing antigen
10 recognized by the antiserum. After a final washing, the filter was developed in a substrate medium containing 33 μ l NBT (50 mg/ml stock solution maintained at 5°C) mixed with 16 μ l BCIP (50 mg/ml stock solution maintained at 5°C) in 5 ml of alkaline
15 phosphatase buffer (100 mM Tris, 9.5, 100 mM NaCl, 5 mM $MgCl_2$). Purple color appeared at points of antigen production, as recognized by the antiserum.

D. Screening Plating

20 The areas of antigen production determined in the previous step were replated at about 100-200 pfu on an 82 mm plate. The above steps, beginning with a 5-8 hour incubation, through NBT-BCIP development, were repeated in order to plaque purify phage secreting an
25 antigen capable of reacting with the ET-NANB antibody. The identified plaques were picked and eluted in phage buffer (Maniatis, p. 443).

E. Epitope Identification

30 A series of subclones derived from the original PTZ-KF1 (ET1.1) plasmid from Example 2 were isolated using the same techniques described above. Each of these five subclones were immunoreactive with a pool of anti-ET antisera noted in C. The subclones contained
35 short sequences from the "reverse" sequence set forth previously. The beginning and ending points of the sequences in the subclones (relative to the full

"reverse" sequence), are identified in the table below.

5

TABLE 1

	<u>Subclone</u>	<u>Position in "Reverse" Sequence</u>	
		<u>5'-end</u>	<u>3'-end</u>
	Y1	522	643
10	Y2	594	667
	Y3	508	665
	Y4	558	752
	Y5	545	665

15

Since all of the gene sequences identified in the table must contain the coding sequence for the epitope, it is apparent that the coding sequence for the epitope falls in the region between nucleotide 594 (5'-end) and 643 (3'-end). Genetic sequences equivalent to and complementary to this relatively short sequence are therefore particularly preferred aspects of the present invention, as are peptides produced using this coding region.

20

25

A second series of clones identifying an altogether different epitope was isolated with only Mexican serum.

30

TABLE 2

	<u>Subclone</u>	<u>Position in "Forward" Sequence</u>	
		<u>5'end</u>	<u>3' end</u>
	ET 2-2	2	193
	ET 8-3	2	135
35	ET 9-1	2	109
	ET 13-1	2	101

The coding system for this epitope falls between nucleotide 2 (5'-end) and 101 (3'-end). Genetic sequences related to this short sequence are therefore also preferred, as are peptides produced using this coding region.

While the invention has been described with reference to particular embodiments, methods, construction and use, it will be apparent to those skilled in the art that various changes and modifications can be made without departing from the invention.

15

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WE CLAIM:

1. A protein derived from an enterically transmitted non-A/non-B viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4 and having ATCC deposit no. 67717.
2. The protein of claim 1, which is encoded by a coding region within said 1.33 kb EcoRI insert.
3. A recombinant protein derived from an enterically transmitted nonA/nonB viral hepatitis agent whose genome contains a region which is homologous to a duplex DNA having a first sequence:

1	11	21	31	41	51
*	*	*	*	*	*
20	CGAATTCGCCCAACTGATGGAAGGCACTAATCTGGCAAGACCTGTCCCTGTTGCAGCTGT				
61	71	81	91	101	111
*	*	*	*	*	*
25	TCTACCACCCTGCCCCGAGCTCGAACAGGGCCTTCTCTACCTGCCCCAGGAGCTCACCAC				
121	131	141	151	161	171
*	*	*	*	*	*
30	CTGTGATAGTGTGTAACATTTGAATTAACAGACATTGTGCACTGCCGCATGGCCGCCCC				

52

181 191 201 211 221 231
* * * * * *
GAGCCAGCGCAAGGCCGTGCTGTCCACACTCGTGGGCCGCTACGGCGTCGCACAAAGCTC

5 241 251 261 271 281 291
* * * * * *
TACAATGCTTCCCACTCTGATGTTCCGCGACTCTCTCGCCCGTTTTATCCCGGCCATTGGC

10 301 311 321 331 341 351
* * * * * *
CCCGTACAGGTTACAACCTGTGAATTGTACGAGCTAGTGGAGGCCATGGTTCGAGAAGGGC

15 361 371 381 391 401 411
* * * * * *
CAGGATGGCTCCGCCGTCTTGAGCTTGATCTTTGCAACCGTGACGTGTCCAGGATCACC

20 421 431 441 451 461 471
* * * * * *
TTCTTCCAGAAAGATTGTAACAAGTTCACCACAGGTGAGACCATTGCCCATGGTAAAGTG

25 481 491 501 511 521 531
* * * * * *
GGCCAGGGCATCTCGGCCTGGAGCAAGACCTTCTGCGCCCTCTTTGGCCCTTGGTTCCGC

30 541 551 561 571 581 591
* * * * * *
GCTATTGAGAAGGCTATTCTGGCCCTGCTCCCTCAGGGTGTGTTTTACGGTGATGCCTTT

35 601 611 621 631 641 651
* * * * * *
GATGACACCGTCTTCTCGGCGGCTGTGGCCGAGCAAAGGCATCCATGGTGTGTTGAGAAT

661 671 681 691 701 711
* * * * * *
GACTTTTCTGAGTTTGACTCCACCCAGAATAACTTTTCTCTGGGTCTAGAGTGTGCTATT

721 731 741 751 761 771
* * * * * *
ATGGAGGAGTGTGGGATGCCGCAGTGGCTCATCCGCCTGTATCACCTTATAAGGTCTGCG

5 781 791 801 811 821 831
* * * * * *
TGGATCTTGCAGGCCCCGAAGGAGTCTCTGCGAGGGTTTGGGAAGAAACACTCCGGTGAG

10 841 851 861 871 881 891
* * * * * *
CCCGGCACTCTTCTATGGAATACTGTCTGGAATATGGCCGTTATTACCCACTGTTATGAC

15 901 911 921 931 941 951
* * * * * *
TTCCGCGATTTTCAGGTGGCTGCCCTTAAAGGTGATGATTCGATAGTGCTTTGCAGTGAG

20 961 971 981 991 1001 1011
* * * * * *
TATCGTCAGAGTCCAGGAGCTGCTGTCCTGATCGCCGGCTGTGGCTTGAAGTTGAAGGTA

25 1021 1031 1041 1051 1061 1071
* * * * * *
GATTTCCGCCCCGATCGGTTTGTATGCAGGTGTTGTGGTGGCCCCCGGCCTTGGCGCGCTC

30 1081 1091 1101 1111 1121 1131
* * * * * *
CCTGATGTTGTGCGCTTCGCCGGCCGGCTTACCGAGAAGAATTGGGGCCCTGGCCCTGAG

35 1141 1151 1161 1171 1181 1191
* * * * * *
CGGGCGGAGCAGCTCCGCCTCGCTGTTAGTGATTTCTCCGCAAGCTCACGAATGTAGCT

1201 1211 1221 1231 1241 1251
* * * * * *
CAGATGTGTGTGGATGTTGTTTCCCGTGTTTATGGGGTTTCCCCTGGACTCGTTTCATAAC

1261 1271 1281 1291 1301 1311
* * * * *
CTGATTGGCATGCTACAGGCTGTTGCTGATGGCAAGGCACATTCTACTGAGTCAGTAAAA

5 1321 1331 1341
* * *
CCAGTGCTCGACCGGAATTCGAGC

or a second sequence

10 1 11 21 31 41 51
* * * * *
GCTCGAATTCGGTCGAGCACTGGTTTTACTGACTCAGTGAAATGTGCCTTGCCATCAGC

15 61 71 81 91 101 111
* * * * *
AACAGCCTGTAGCATGCCAATCAGGTTATGAACGAGTCCAGGGGAAACCCATAAACACG

20 121 131 141 151 161 171
* * * * *
GGAAACAACATCCACACACATCTGAGCTACATTCGTGAGCTTCCGGAGGAAATCACTAAC

25 181 191 201 211 221 231
* * * * *
AGCGAGGCGGAGCTGCTCCGCCCCTCAGGGCCAGGGCCCCAATTCTTCTCGGTAAGCCG

30 241 251 261 271 281 291
* * * * *
GCCGCGGAAGCGCACAAATCAGGGAGCGCGCCAAGGCCGGGGGCCACCACAACACCTGC

35 301 311 321 331 341 351
* * * * *
ATACAAACCGATCGGGCGGAAATCTACCTTCAACTTCAAGCCACAGCCGGCGATCAGGAC

40 361 371 381 391 401 411
* * * * *
AGCAGCTCCTGGACTCTGACGATACTCACTGCAAAGCACTATCGAATCATCACCTTTAAA

421 431 441 451 461 471
* * * * *
GGCAGCCACCTGAAAATCGCGGAAGTCATAACAGTGGGTAATAACGGCCATATTCCAGAC
5
481 491 501 511 521 531
* * * * *
AGTATTCCATAGAAGAGTGCCGGGCTCACGGAGTGTTCCTCCAAAACCCCTCGCAGAGA
10 541 551 561 571 581 591
* * * * *
CTCCTTCGGGGCCTGCAAGATCCACGCAGACCTTATAAGGTGATACAGGCGGATGAGCCA
601 611 621 631 641 651
15 * * * * *
CTGCGGCATCCCACACTCCTCCATAATAGCACACTCTAGACCCAGAGAAAAGTTATTCTG
661 671 681 691 701 711
* * * * *
20 GGTGGAGTCAAACCTCAGAAAAGTCATTCTCAAACACCATGGATGCCTTTGCTGCGGCCAC
721 731 741 751 761 771
* * * * *
25 AGCCGCCGAGAAGACGGTGTCAATAGGCATCACCGTAAACACACCCTGAGGGAGCAG
781 791 801 811 821 831
* * * * *
GGCCAGAATAGCCTTCTCAATAGCGCGGAACCAAGGGCCAAAGAGGGCGCAGAAGGTCTT
30 841 851 861 871 881 891
* * * * *
GCTCCAGGCCGAGATGCCCTGGCCCACTTTACCATGGGCAATGGTCTCACCTGTGGTGAA
901 911 921 931 941 951
35 * * * * *
CTTGTTACAATCTTTCTGGAAGAAGGTGATCCTGGACACGTCACGGTTGCAAAGATCAAG

961 971 981 991 1001 1011
 * * * * * *
 CTC AAGGACGGCGGAGCCATCCTGGCCCTTCTCGACCATGGCCTCCACTAGCTCGTACAA
 5 1021 1031 1041 1051 1061 1071
 * * * * * *
 TTCACAAGTTGTAACTGTACGGGGCCAATGGCCGGGATAAAACGGGCGAGAGAGTCGCG
 1081 1091 1101 1111 1121 1131
 10 * * * * * *
 AACATCAGAGTGGGAAGCATTGTAGAGCTTTGTGCGACGCCGTAGCGGCCACGAGTGTG
 1141 1151 1161 1171 1181 1191
 * * * * * *
 15 GACAGCACGGCCTTGGCTGGCTCGGGGCGGCCATGCGGCAGTGCACAATGTCTGTTAAT
 1201 1211 1221 1231 1241 1251
 * * * * * *
 TCAAATGTTACGACACTATCACAGGTGGTGAGCTCCTGGGGCAGGTAGAGAAGGCCCTGT
 20 1261 1271 1281 1291 1301 1311
 * * * * * *
 TCGAGCTCGGGGCAGGGTGGTAGAACAGCTGCAACAGGGACAGGTCTTGCCAGATTAGTG
 1321 1331 1341
 25 * * *
 CCTTCATCAGTTGGCGGAATTCG.

- 30 4. A protein which is (a) immunoreactive with antibodies
 present in individuals infected with enterically
 transmitted nonA/nonB and (b) derived from a viral
 hepatitis agent whose genome contains a region which is
 homologous to the 1.33 kb DNA EcoRI insert present in
 plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4,
 35 and having ATCC deposit no. 67717.

5. The protein of claim 4, which is encoded by a coding region within said 1.33 kb EcoRI insert.
6. A method of detecting infection by enterically transmitted nonA/nonB hepatitis viral agent in a test individual, comprising:
- 5 providing a peptide antigen which is (a) immunoreactive with antibodies present in individuals infected with enterically transmitted nonA/nonB hepatitis and (b) derived from a viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4, and having ATCC deposit no. 67717,
- 10 reacting serum from the test individual with such antigen, and examining the antigen for the presence of bound antibody.
7. The method of claim 6, wherein the serum antibody is an IgM or IgG antibody, or a mixture of both, the antigen provided is attached to a support, said reacting includes contacting such serum with the support and said examining includes reacting the support and bound serum antibody with a reporter-labeled anti-human antibody.
- 20 7. The method of claim 6, wherein the serum antibody is an IgM or IgG antibody, or a mixture of both, the antigen provided is attached to a support, said reacting includes contacting such serum with the support and said examining includes reacting the support and bound serum antibody with a reporter-labeled anti-human antibody.
- 25 8. A kit for ascertaining the presence of serum antibodies which are diagnostic of enterically transmitted nonA/nonB hepatitis infection, comprising
- 30 a support with surface-bound recombinant peptide antigen which is (a) immunoreactive with antibodies present in individuals infected with enterically transmitted nonA/nonB viral hepatitis agent and (b) derived from a viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4, and having ATCC deposit no.
- 35

67717,

a reporter-labeled anti-human antibody.

9. A DNA fragment derived from an enterically transmitted nonA/nonB viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4 and having ATCC deposit no. 67717.
10. The fragment of claim 9, which is derived from said 1.33 kb EcoRI insert.

11. A DNA fragment derived from an enterically transmitted nonA/nonB viral hepatitis agent whose genome contains a region which is homologous to a duplex DNA fragment within a first sequence:

1	11	21	31	41	51
*	*	*	*	*	*
CGAATTCCGCCAACTGATGGAAGGCACTAATCTGGCAAGACCTGTCCCTGTTGCAGCTGT					
20	61	71	81	91	101
	*	*	*	*	*
TCTACCACCCTGCCCCGAGCTCGAACAGGGCCTTCTCTACCTGCCCCAGGAGCTCACCAC					
25	121	131	141	151	161
	*	*	*	*	*
CTGTGATAGTGTGTAACATTTGAATTAACAGACATTGTGCACTGCCGCATGGCCGCCCC					
30	181	191	201	211	221
	*	*	*	*	*
GAGCCAGCGCAAGGCCGTGCTGTCCACACTCGTGGGCCGCTACGGCGTCGCACAAAGCTC					
	241	251	261	271	281
	*	*	*	*	*
35	TACAATGCTTCCCACTCTGATGTTTCGCGACTCTCTCGCCCGTTTATCCCGGCCATTGGC				

301 311 321 331 341 351
* * * * *
CCCGTACAGGTTACAACCTTGTGAATTGTACGAGCTAGTGGAGGCCATGGTTCGAGAAGGGC
5
361 371 381 391 401 411
* * * * *
CAGGATGGCTCCGCCGTCCTTGAGCTTGATCTTTGCAACCGTGACGTGTCCAGGATCACC
10
421 431 441 451 461 471
* * * * *
TTCTTCCAGAAAGATTGTAACAAGTTCACCACAGGTGAGACCATTGCCCATGGTAAAGTG
15
481 491 501 511 521 531
* * * * *
GGCCAGGGCATCTCGGCCTGGAGCAAGACCTTCTGCGCCCTCTTTGGCCCTTGTTCCGC
20
541 551 561 571 581 591
* * * * *
GCTATTGAGAAGGCTATTCTGGCCCTGCTCCCTCAGGGTGTGTTTACGGTGATGCCTTT
25
601 611 621 631 641 651
* * * * *
GATGACACCGTCTTCTCGGCGGCTGTGGCCGCAGCAAAGGCATCCATGGTGTTTGAGAAT
30
661 671 681 691 701 711
* * * * *
GACTTTTCTGAGTTTGA TCCACCCAGAATAACTTTTCTCTGGGTCTAGAGTGTGCTATT
35
721 731 741 751 761 771
* * * * *
ATGGAGGAGTGTGGGATGCCGCAGTGGCTCATCCGCCTGTATCACCTTATAAGGTCTGCG
781 791 801 811 821 831
* * * * *
TGGATCTTGCAGGCCCCGAAGGAGTCTCTGCGAGGGTTTTGGAAGAAACACTCCGGTGAG

841 851 861 871 881 891
* * * * *
CCCGGCACTCTTCTATGGAATACTGTCTGGAATATGGCCGTTATTACCCACTGTTATGAC

5 901 911 921 931 941 951
* * * * *
TTCCGCGATTTTCAGGTGGCTGCCCTTTAAAGGTGATGATTCGATAGTGCTTTGCAGTGAG

10 961 971 981 991 1001 1011
* * * * *
TATCGTCAGAGTCCAGGAGCTGCTGTCCTGATCGCCGGCTGTGGCTTGAAGTTGAAGGTA

15 1021 1031 1041 1051 1061 1071
* * * * *
GATTTCCGCCCCGATCGGTTTGTATGCAGGTGTTGTGGTGGCCCCCGGCCTTGGCGCGCTC

20 1081 1091 1101 1111 1121 1131
* * * * *
CCTGATGTTGTGCGCTTCGCCGGCCGGCTTACCGAGAAGAATTGGGGCCCTGGCCCTGAG

1141 1151 1161 1171 1181 1191
* * * * *
CGGGCGGAGCAGCTCCGCCTCGCTGTTAGTGATTTCTCCGCAAGCTCACGAATGTAGCT

25 1201 1211 1221 1231 1241 1251
* * * * *
CAGATGTGTGTGGATGTTGTTTCCCGTGTTTATGGGGTTTCCCCTGGACTCGTTCATAAC

30 1261 1271 1281 1291 1301 1311
* * * * *
CTGATTGGCATGCTACAGGCTGTTGCTGATGGCAAGGCACATTTCACTGAGTCAGTAAAA

1321 1331 1341
* * *
35 CCAGTGCTCGACCGGAATTCGAGC

or a second sequence

1 11 21 31 41 51
* * * * * *
GCTCGAATTCGGTTCGAGCACTGGTTTTACTGACTCAGTGAAATGTGCCTTGCCATCAGC
5
61 71 81 91 101 111
* * * * * *
AACAGCCTGTAGCATGCCAATCAGGTTATGAACGAGTCCAGGGGAAACCCATAAACACG
10
121 131 141 151 161 171
* * * * * *
GGAAACAACATCCACACACATCTGAGCTACATTCGTGAGCTTGCGGAGGAAATCACTAAC
15
181 191 201 211 221 231
* * * * * *
AGCGAGGCGGAGCTGCTCCGCCCCTCAGGGCCAGGGCCCCAATTCTTCTCGGTAAGCCG
20
241 251 261 271 281 291
* * * * * *
GCCGGCGAAGCGCACAAACATCAGGGAGCGCGCCAAGGCCGGGGGCCACCACAACACCTGC
25
301 311 321 331 341 351
* * * * * *
ATACAAACCGATCGGGCGGAAATCTACCTTCAACTTCAAGCCACAGCCGGCGATCAGGAC
30
361 371 381 391 401 411
* * * * * *
AGCAGCTCCTGGACTCTGACGATACTCACTGCAAAGCACTATCGAATCATCACCTTTAAA
35
421 431 441 451 461 471
* * * * * *
GGCAGCCACCTGAAAATCGCGGAAGTCATAACAGTGGGTAATAACGGCCATATTCCAGAC
481 491 501 511 521 531
* * * * * *
AGTATTCCATAGAAGAGTGCCGGGCTCACCAGAGTGTTTCTTCCAAAACCTCGCAGAGA

541 551 561 571 581 591
* * * * * *
CTCCTTCGGGGCCTGCAAGATCCACGCAGACCTTATAAGGTGATACAGGCGGATGAGCCA

5 601 611 621 631 641 651
* * * * * *
CTGCGGCATCCCACACTCCTCCATAATAGCACACTCTAGACCCAGAGAAAAGTTATTCTG

10 661 671 681 691 701 711
* * * * * *
GGTGGAGTCAAACCTCAGAAAAGTCATTCTCAAACACCATGGATGCCTTTGCTGCGGCCAC

15 721 731 741 751 761 771
* * * * * *
AGCCGCCGAGAAGACGGTGTCTATCAAAGGCATCACCGTAAACACACCCTGAGGGAGCAG

20 781 791 801 811 821 831
* * * * * *
GGCCAGAATAGCCTTCTCAATAGCGCGGAACCAAGGGCCAAAGAGGGCGCAGAAGGTCTT

25 841 851 861 871 881 891
* * * * * *
GCTCCAGGCCGAGATGCCCTGGCCACTTTACCATGGGCAATGGTCTCACCTGTGGTGAA

30 901 911 921 931 941 951
* * * * * *
CTTGTTACAATCTTTCTGGAAGAAGGTGATCCTGGACACGTCACGGTTGCAAAGATCAAG

35 961 971 981 991 1001 1011
* * * * * *
CTCAAGGACGGCGGAGCCATCCTGGCCCTTCTCGACCATGGCCTCCACTAGCTCGTACAA

1021 1031 1041 1051 1061 1071
* * * * * *
TTCACAAGTTGTAACCTGTACGGGGCCAATGGCCGGGATAAAACGGGCGAGAGAGTCGCG


```

1081      1091      1101      1111      1121      1131
*          *          *          *          *          *
AACATCAGAGTGGGAAGCATTGTAGAGCTTTGTGCGACGCCGTAGCGGCCCCACGAGTGTG

5  1141      1151      1161      1171      1181      1191
*          *          *          *          *          *
GACAGCACGGCCTTGCGCTGGCTCGGGGCGGCCATGCGGCAGTGACACAATGTCTGTTAAT

10 1201      1211      1221      1231      1241      1251
*          *          *          *          *          *
TCAAATGTTACGACACTATCACAGGTGGTGAGCTCCTGGGGCAGGTAGAGAAGGCCCTGT

1261      1271      1281      1291      1301      1311
*          *          *          *          *          *
15 TCGAGCTCGGGGCAGGGTGGTAGAACAGCTGCAACAGGGACAGGTCTTGCCAGATTAGTG

1321      1331      1341
*          *          *
CCTTCCATCAGTTGGCGGAATTCG.

```

20

12. The DNA fragment of claim 11, wherein said fragment contains a coding sequence homologous to nucleotides 2 through 101 of said first sequence, nucleotides 594 through 643 of said second sequence, or a sequence complementary to said coding sequences.

25

13. A kit comprising, in a container or separate containers, a pair of single-strand primers derived from non-homologous regions of opposite strands of a DNA duplex fragment derived from an enterically transmitted viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4 and having ATCC deposit no. 67717.

30

14. The kit of claim 13, which are derived from opposite strands of the EcoRI duplex insert in said plasmid.

35

15. A method for detecting the presence of an enterically transmitted nonA/nonB hepatitis viral agent in a biological sample, comprising
- 5 preparing a mixture of duplex DNA fragments derived from the sample, denaturing the duplex fragments, adding to the denatured DNA fragments, a pair of single-strand primers derived from non-
- 10 homologous regions of opposite strands of a DNA duplex fragment derived from an enterically transmitted viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4,
- 15 and having ATCC deposit no. 67717, hybridizing said primers to homologous-sequence region of opposite strands of such duplex DNA fragments derived from enterically transmitted nonA/nonB hepatitis agent,
- 20 reacting the primed fragment strands with DNA polymerase in the presence of DNA nucleotides, to form new DNA duplexes containing the primer sequences, and repeating said denaturing, adding, hybridizing and reacting steps, until a desired degree of
- 25 amplification of sequences is achieved.
16. The method of claim 15, wherein the primers are derived from opposite strands the EcoRI duplex insert in said plasmid.
- 30 17. The method of claim 15, for detecting the presence of viral agent in a sample of cultured cells infected with the agent.
- 35 18. A vaccine for immunizing an individual against enterically transmitted nonA/nonB hepatitis viral agent comprising, in a pharmacologically acceptable adjuvant,

a recombinant protein derived from an enterically transmitted nonA/nonB viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4, and having ATCC deposit no. 67717.

19. The vaccine of claim 18, wherein the protein is derived from the EcoRI insert in said plasmid.
20. In a method of isolating an enterically transmitted non-A/nonB viral agent or a nucleic acid fragment produced by the agent, an improvement which comprises:
utilizing, as a source of said agent, bile obtained from a human or cynomolgus monkey having an active infection of enterically transmitted non-A/non-B hepatitis.
21. The method of claim 20, wherein the bile is obtained from an infected cynomologus monkey.
22. Human polyclonal anti-serum obtained from a human immunized with a protein derived from an enterically transmitted non-A/non-B viral hepatitis agent whose genome contains a region which is homologous to the 1.33 kb DNA EcoRI insert present in plasmid pTZ-KF1(ET1.1) carried in E. coli strain BB4 and having ATCC deposit no. 67717.

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Fig. 1

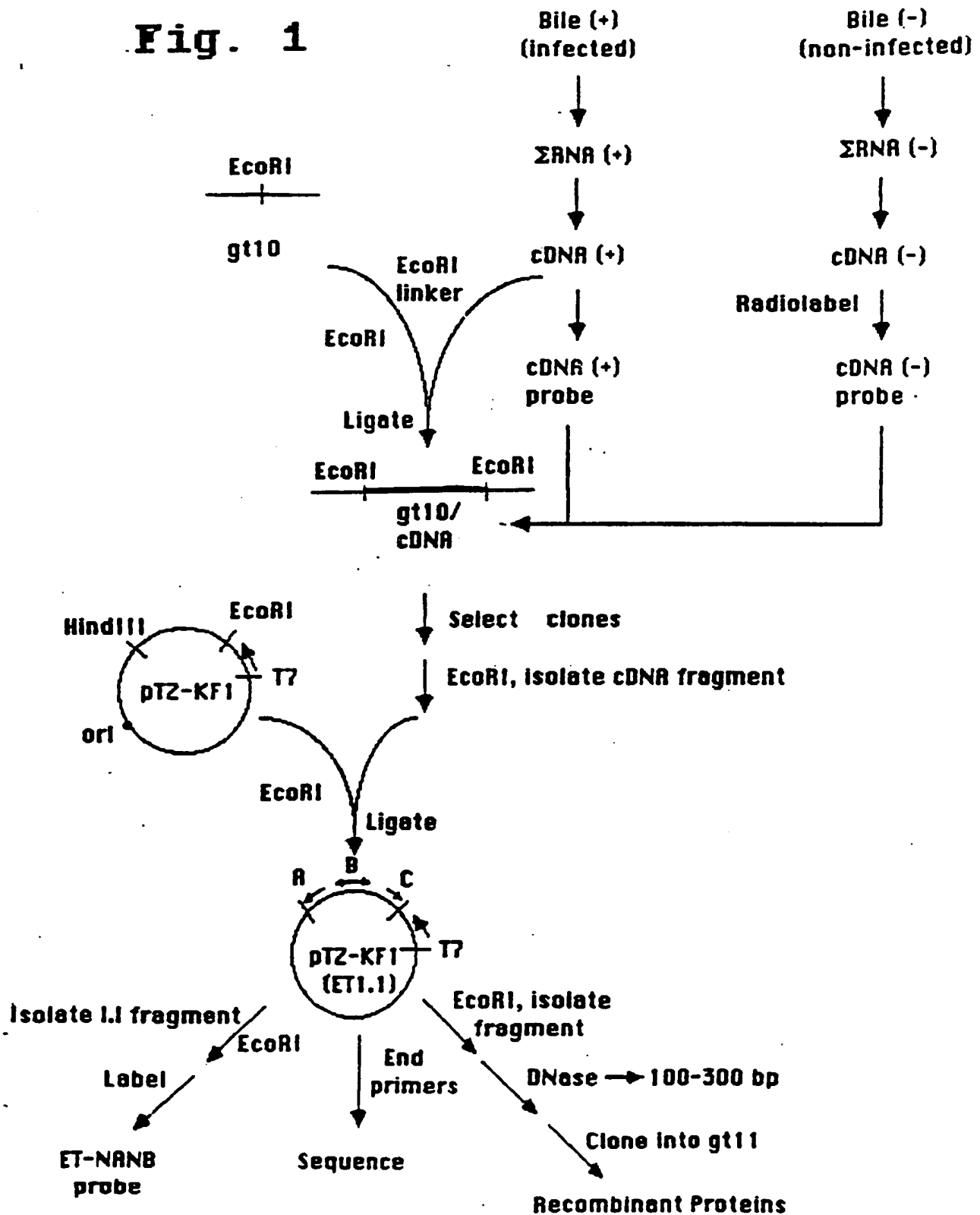
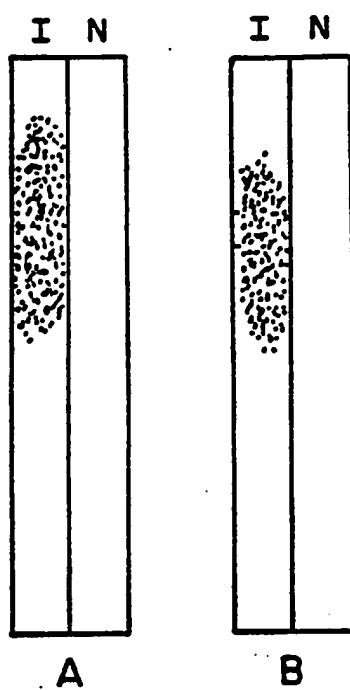


FIG. 2



INTERNATIONAL SEARCH REPORT

International Application No. PCT/US89/02435

I. CLASSIFICATION OF SUBJECT MATTER * See also classification symbols appearing on page 1
According to International Patent Classification (IPC) or to both National Classification and IPC U.S. CL. 435/5,810
436/513,518,536,543,820;530/324,350,806,808,826;536/27 INT.CL.4
C07H 17/00; C07K 7/10, 15/04; C12Q 1/70;GOIN 33/531,536,543,576

II. FIELDS SEARCHED

Minimum Documentation Searched *

Classification System

Classification Symbols

US. CL.

435/5,810; 436/513,518,536,543,820
530/324,350,806,808,826; 536/27

Documentation Searched other than Minimum Documentation
to the Extent that such Documents are Included in the Fields Searched *

BIOSIS SEARCH: (ET or ENTERIC?) (5A) (NANB? OR NON(W)B)

III. DOCUMENTS CONSIDERED TO BE RELEVANT *

Category *	Citation of Document, ** with indication, where appropriate, of the relevant passages †	Relevant to Claim No. ‡
Y	WO,A, 85/01517 (MASSACHUSETTS INSTITUTE OF TECHNOLOGY) 30 September 1983. See page 5, line 6-page 6, line 22; page 14, lines 10-27.	1-12
Y	US,A 4,591,552 (NEURATH) 29 September 1982. See col. 15, lines 22-29; col. 19, lines 58-68.	6-8
Y	The Lancet, vol 1, no. 8585, issued 12 March 1988, V.A. ARANKALLE ET AL, "Aetiological associated of a virus like particle with enterically transmitted non-A, non-B hepatitis", pages 550-554. See Summary.	1-12

* Special categories of cited documents: †

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"A" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search

Date of Mailing of this International Search Report

03 August 1989

19 OCT 1989

International Searching Authority

Signature of Authorized Officer

ISA/US

David A. Saunders

III DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category	Citation of Document, with indication where appropriate, of the relevant passages	Relevant to Claim No
Y	<u>Journal of General Virology</u> , vol. 69, no.3 issued March 1988, D. BRADLEY ET AL., "Aetiological agent of enterically transmitted non-A non-B hepatitis", pages 731-738, See Summary.	1-12
Y	<u>Proceedings of the National Academy of Sciences USA</u> , vol. 84 no.17, issued September 1987, D.W. BRADLEY ET AL, "Enterically transmitted non-A, non-B hepatitis. Serial passage of disease in cynomolgus macaques and tamarins and recovery of disease-associated 27-to 34-NM viruslike particles", pages 6277-6281. See abstract.	1-12
A,P	<u>The Journal of Infectious Disease</u> , vol. 159, no.6, issued June 1989, K. KRAWCZYNSKI ET AL, "Enterically transmitted non-A, non-B, hepatitis: Identification of virus-associated antigen in experimentally infected cynomolgus macques", pages 1042-1049. See page 1042 first full paragraph; page 1048 second full paragraph.	6

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

V ☐ OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE¹

This international search report has not been established in respect of certain claims under Article 17(2) (a) for the following reasons:

1. ☐ Claim numbers _____, because they relate to subject matter¹² not required to be searched by this Authority, namely:

2. ☐ Claim numbers _____, because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out¹³, specifically:

3. ☐ Claim numbers _____, because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 6.4(a).

VI. ☒ OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING³

This International Searching Authority found multiple inventions in this international application as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims of the international application.
2. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims of the international application for which fees were paid, specifically claims:
3. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:
4. ☐ As all searchable claims could be searched without effort justifying an additional fee, the International Searching Authority did not invite payment of any additional fee.

Remark on Protest

- ☐ The additional search fees were accompanied by applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.